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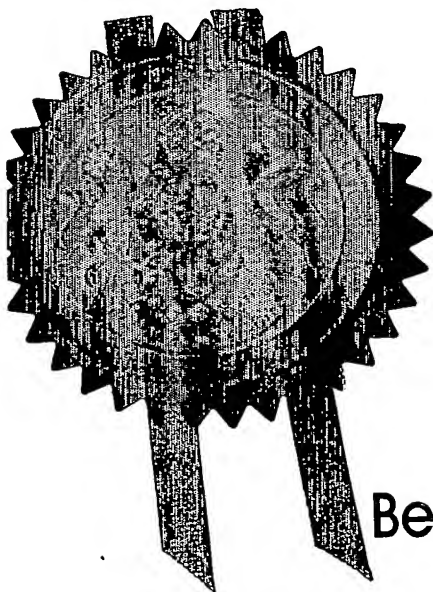
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26 APR 2002

1. Your reference	300367GB/JND/JE/ms	29APR02 E714442-1 D00068
2. Patent application number (The Patent Office will fill in this part)	0209652.7	
3. Full name, address and postcode of the or of each applicant (underline all surnames)	Cambridge University Technical Services Ltd. The Old Schools Trinity Lane Cambridge CB2 1TS United Kingdom	
Patents ADP number (if you know it)	6956809004	
If the applicant is a corporate body, give the country/state of its incorporation	United Kingdom	

Title of the invention	Solution-Processable Phosphorescent Materials
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Name of your agent (if you have one)	Page White & Farrer
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"Address for service" in the United Kingdom to which all correspondence should be sent (including the postcode)	54 Doughty Street London WC1N 2LS United Kingdom
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Patents ADP number (if you know it)	1255003 ✓
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Is a statement of inventorship and of right to grant of a patent required in support of this request? (Answer 'Yes' if: a) any applicant named in part 3 is not an inventor, or b) there is an inventor who is not named as an applicant, or c) any named applicant is a corporate body See note (d))	Yes
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Drawing(s) 12 + 12 - JMC

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12. Name and daytime telephone number of person to contact in the United Kingdom

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SOLUTION-PROCESSABLE PHOSPHORESCENT MATERIALS

The present invention relates to novel materials that are capable of luminescence; to a method of making the novel materials and to uses of the novel materials.

Luminescent conjugated polymers are a new technologically important class of materials that will be used in light emitting display devices for the next generation of information technology based consumer products. The principle interest in the use of polymers, as opposed to inorganic semiconducting and organic dye materials, lies in the scope for low-cost manufacturing, using solution-processing of film-forming materials. Since the last decade much effort has been devoted to the improvement of the emission efficiency of organic light emitting diodes (OLED's) either by developing highly efficient materials or efficient device structures.

In OLED's, electrons and holes are injected from opposite electrodes and are combined to form two types of excitons; spin-symmetric triplets and spin-antisymmetric singlets in a theoretical ratio of 3:1. Radiative decay from the singlets is fast (fluorescence), but from the triplets (phosphorescence) it is formally forbidden by the requirement of the spin conservation.

Initially spurred on by this understanding that the maximum internal quantum efficiency of an OLED was limited to 25% the idea of transferring both singlets and triplets to a phosphorescent dopant was conceived. Such a *phosphor* ideally is able to accept both singlet and triplet excitons from the organic material and generate luminescence, particularly electroluminescence from both.

This idea is still highly applicable even given recent studies questioning the 3:1 triplet to singlet ratio predicted by the spin-independent recombination model. Recent studies suggest that the proportion of triplet excitons generated in small molecule devices is indeed close to 75%, (Baldo, M.A.; O'Brien, D.F.; Thompson, M.E.; Forrest, S.R. *Phys. Rev. B* 1999, 60, 14422) whereas in an electrically excited conjugated polymer it is suggested to be around 50%, (Cao, Y.; Parker, I.D.; Yu, G.; Zhang, C.; Heeger, A.J. *Nature* 1999, 397, 414 and Wilson, J.S.; Dhoot, A.S.; Seeley, A.J.A.B.; Khan, M.S.; Köhler, A.; Friend, R.H. *Nature* 2001, 413, 828). Evidence for

the copious generation of triplet excitons in polymer LED's has been obtained by magnetic and optical observations.

In the past few years many have studied the incorporation by blending of phosphorescent materials into a semiconductive layer. Good results have been achieved for OLED's based on small molecule blends incorporating a phosphorescent dopant. Lanthanide metal chelates are by far the most common triplet emitters used in OLEDs. A paper notable for this work reports a device utilising a blend of $\text{Eu}(\text{dnm})_3\text{phen}$ in CN-PPP with a quantum efficiency of 1.1%. [*Adv. Mater.*, 1999, 11, 1349.]. To date, equivalently doped conjugated polymers have not been successful.

One of the reasons for the lack of success with polymers is that blends of polymers are sensitive to the concentration of the triplet harvester guest in the polymer host. Therefore even at relatively low concentrations the guest can phase separate leading to morphology changes such as aggregation and problems with light scattering, electrochemical breakdown and luminescence quenching.

In view of the aforementioned problems, predictions for achieving good efficiency with phosphorescent, particularly electrophosphorescent, polymer OLED's have been pessimistic. As such, the art has focused on achieving optimal singlet emission (fluorescence) from polymer OLED's.

In this regard, it is to be noted that luminescent polymers containing metal complexes are known in this art. However, in keeping with the above, these materials act predominantly as fluorescent materials and not phosphorescent materials. An example of such a material is given in *Macromol. Rapid. Commun.* 18, 1009-1016 (1997).

In view of the above, it will be appreciated that there still remains a need to provide polymer OLED's having good efficiency.

Accordingly, it is an aim of the present invention to at least partially address this need by providing a new emissive material for use in an OLED.

It is a further aim of the present invention to provide methods for making the new emissive material.

It is still a further aim of the present invention to provide uses for the new emissive material.

Accordingly, the present invention provides a material capable of luminescence comprising a polymer or oligomer; and an organometallic characterised in that the organometallic is covalently bound to the polymer or oligomer and the nature, location and/or proportion of the polymer or oligomer and of the organometallic in the material are selected so that the luminescence predominantly is phosphorescence.

The present inventors have addressed the aforementioned need for polymer OLED's having good efficiency by providing a new material that is capable predominantly of phosphorescence, preferably electrophosphorescence. In the present invention, by "phosphorescence" is meant luminescence by radiative decay from a triplet energy level.

The present inventors have addressed at least partially the aforementioned need by harvesting previously wasted triplet excitons. This is completely opposed to the currently favoured work concentrating on optimising harvest of singlet excitons from polymers.

The present material generally is superior to a polymer blend incorporating a phosphorescent dopant. This is because problems relating to morphology changes such as aggregation and phase separation are avoided with the present material. The controlled structure of the present material means that the location and mobility of the organometallic in the material is spatially controlled. This spatial control enables control of the interaction between the polymer or oligomer and the organometallic. This, in turn, enables manipulation, particularly to some extent of the energy levels of the material in its excited state when the organometallic is conjugatively linked to the polymer or oligomer because of synergistic effects between the polymer or oligomer and the organometallic.

The present materials are advantageous in that they give bright phosphorescence in solution-processed devices. Also, recent reports suggesting the triplet states of conjugated polymers such as poly(3-alkylthiophene) [*Macromol.*, 1993, 26, 2954.] and PPV derivatives [*Chem. Mater.*, 1999, 11, 1302.] can sensitise the generation of singlet oxygen, which can degrade the polymer.

Also, advantageously, the present material usually is homogenous and solution-processable.

The nature, location and/or proportion of the polymer or oligomer and of the organometallic in the material is selected so that the luminescence predominantly is phosphorescence. Preferably, greater than 70% of the luminescence is phosphorescence, even more preferably, greater than 90% of the luminescence is phosphorescence. Most preferably, 95 to 100% of the luminescence is phosphorescence.

The nature, location and/or proportion of the polymer or oligomer and of the organometallic in the material should be selected so that the energy levels of the polymer or oligomer and organometallic are complimentary to one another. This will allow the transfer of both singlets and triplets from the polymer or oligomer in its excited state to the organometallic. Theoretically, 100% transfer of both singlets and triplets from the polymer or oligomer can be achieved. Predominantly phosphorescent emission then will occur. In order for this to occur, generally but not essentially, the singlet and triplet energy levels of the organometallic should be lower than the corresponding singlet and triplet energy level of the polymer or oligomer. (Adachi et al. *Appl. Phys. Lett.*, 2001, 79, 2082).

When the organometallic is conjugatively linked to the polymer or oligomer, this may lead to mixing of the energy levels.

Generally, an increase in the extent of conjugation in a material can be said to lower the triplet energy level of that material. As such, controlling the extent of conjugation in the present material can be a useful method of controlling at least partially the triplet energy levels. In this regard, in one embodiment, it is preferred that the

backbone of the polymer or oligomer is partially conjugated, more preferably substantially or even fully conjugated in order to control to some extent the triplet energy level of the polymer or oligomer.

In order to increase conjugation of the polymer or oligomer, it is preferred that the polymer or oligomer comprises an aryl or heteroaryl repeat unit, for example a fluorene or phenyl group, such as a phenylene vinylene group. Thus, in one embodiment, the present material preferably comprises a polyfluorene or polyphenylene.

Preferred aryl or heteroaryl repeat units comprise a group selected from the group consisting of 2,7-linked 9,9 disubstituted fluorene, a p-linked dialkyl phenylene, a p-linked disubstituted phenylene, a phenylene vinylene, a 2,5-linked benzothiadiazole, a 2,5-linked substituted benzothiadiazole, a 2,5-linked disubstituted benzothiadiazole, a 2,5-linked substituted or unsubstituted thiophene or a triarylamine.

Any polymer or oligomer can be used so long as its energy levels are complimentary to the organometallic so that the material is capable predominantly of phosphorescence, preferably electrophosphorescence. To this end, in one embodiment, the present inventors have found that the polymer or oligomer preferably is conducting or semiconducting and, more preferably, semiconducting. In this embodiment, preferably, the polymer or oligomer is capable predominantly of fluorescence (i.e. singlet emission), preferably electrofluorescence, in the absence of the organometallic.

It may be preferable in some embodiments for the polymer or oligomer to include solubilising groups in order to improve the solubility of the material. Preferred solubilising groups include alkyl and alkoxy groups. In this regard, it is preferred that the material is solution-processable.

Usually, the organometallic will be neutral to avoid charge trapping; very stable; and preferably conducting. Importantly the organometallic should be a predominantly triplet emitting material at room temperature.

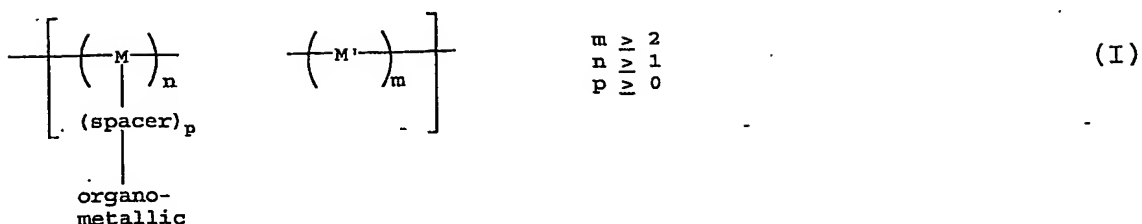
The organometallic will be covalently bonded to the polymer or oligomer in the material. Further, it is preferred that the organometallic is conjugatively bonded to the polymer or oligomer.

Preferably, the organometallic includes a transition metal or lanthanides. The transition metal preferably is a precious metal, more preferably iridium or platinum. It is to be noted that the mechanism for energy transfer will be different where the organometallic is a precious metal as compared with the lanthanide. This difference in mechanism is not important in the present invention so long as the material is capable predominantly of phosphorescence, preferably electrophosphorescence.

It is suggested that any previously known organometallic "phosphor" or a derivative thereof is a good candidate for use as the organometallic in the present invention. In this regard, Platinum, Iridium and Europium complexes are of considerable interest in the present invention.

The nature of the ligands in the organometallic may be selected in order to optimise further compatibility of the organometallic with the polymers or oligomer. To this end, ligands suitably may be selected from the categories: 1) Electron Transport Materials, 2) Hole Transport Materials and 3) (Semi)conducting Materials. In selecting ligands, account should be taken of possible steric hindrance. Typically, materials in these categories preferably comprise a group selected from the group consisting of 2,7-linked 9,9 disubstituted fluorene, a p-linked dialkyl phenylene, a p-linked disubstituted phenylene, a phenylene vinylene, a 2,5-linked benzothiadiazole, a 2,5-linked substituted benzothiadiazole, a 2,5-linked disubstituted benzothiadiazole, a 2,5-linked substituted or unsubstituted thiophene or a triarylamine.

Typically, the present material will contain more than one organometallic. In this case, spatial separation of the organometallics in the material becomes more important. Spatial separation may be controlled by the method of manufacture as described in more detail below or by selecting the organometallic to be pendant from the backbone of the polymer or oligomer. In this regard, the material may have a structure as shown in general formula I below: -



where M and M' each independently comprise an aryl or heteroaryl group.

With a view to further controlling the properties of the material, the spacer group may be used to interrupt conjugation in the material. In this regard alkyl groups and alkyl-ether groups are suitable spacer groups.

The length of the spacer group also can be used to control to some extent the separation of the organometallic from the polymer or oligomer backbone. As such, the spacer group suitably may be a chain of CH₂ groups.

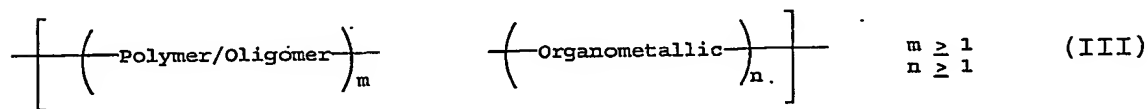
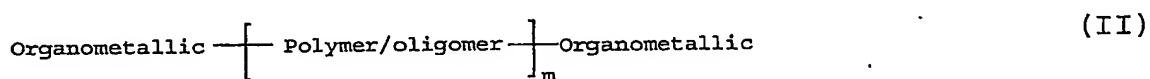
In a preferred material of general formula I, M and M' may be the same or different and each independently comprises a group selected from the group consisting of 2,7-linked 9,9 disubstituted fluorene, a p-linked dialkyl phenylene, a p-linked disubstituted phenylene, a phenylene vinylene, a 2,5-linked benzothiadiazole, a 2,5-linked substituted benzothiadiazole, a 2,5-linked disubstituted benzothiadiazole, a 2,5-linked substituted or unsubstituted thiophene or a triarylamine.

It will be understood that in general formula I, M and M' will be joined to form the backbone of the polymer or oligomer. In this regard, the polymer or oligomer may be a seeded/block or random polymer or oligomer as desired.

One example of a material having general formula I is shown below:

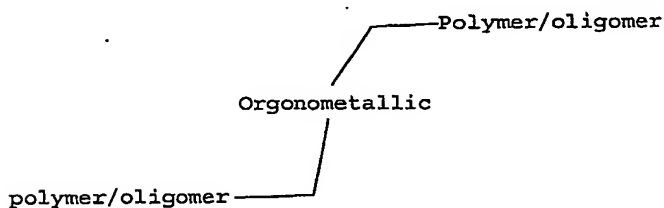
It should be understood that, in general formula I, if it is desirable, one or more additional, preferably aryl or heteroaryl, repeat units that are different to those shown in general formula I may be included in the backbone of the polymer or oligomer.

Where the organometallic is not pendent from the polymer or oligomer backbone, the material may have a structure as shown in general formula II or general formula III below:

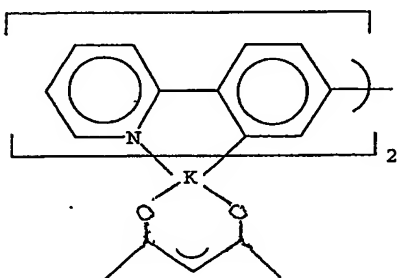


In the materials shown in general formulas II and III, the spatial separation of the organometallics in the material is controlled principally by the method of manufacture.

In general formula III, the polymer/oligomer units and the organometallic units may be linked as desired so that the material is capable predominantly of phosphorescence, preferably electrophosphorescence. One example is shown below:

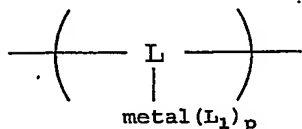


A suitable organometallic for use in the above material is:



In the above, K is a metal, preferably a precious metal, more preferably Ir.

Considering the organometallic as a metal surrounded by an appropriate number of ligands. Usually, in general formulas II and III, the organometallic will be joined to the backbone of the polymer or oligomer through one of its ligands, Ligands around the metal may be the same or different. In this regard, the organometallic repeat unit in the material shown in general formulas II and III may have the structure:



where L is a ligand, preferably a bidentate ligand. Further ligands L_1 (which may be the same or different from one another) are incorporated to satisfy the valency of the

metal. In this regard, L_1 can be a bidentate or monodentate ligand. Preferred L_1 groups include cyclometalating, beta-diketonates. r is chosen accordingly.

Cyclometalated, precious metal complexes such as iridium and platinum complexes are particularly preferred, for example chelate complexes such as platinum polynes, platinum-porphyrins and iridiumtris(phenylpyridine).

The relative proportions of polymer/oligomer and organometallic in the present material must be selected so that the material is capable predominantly of phosphorescence, preferably electrophosphorescence. In this regard, too high a level of organometallic may lead to quenching of the phosphorescence. Alternatively, too low a proportion of organometallic will not receive all of the energy from the polymer or oligomer. Generally, it is preferred that the organometallic is present in the material in an amount in the range of from 0.5 to 70 % by weight, more preferably from 1 to 10 % by weight.

The present materials are extremely useful in OLED's, particularly, for large-area devices like matrix displays. Also, the present materials are of use in electrically pumped organic lasers.

The present materials advantageously can be prepared by the well-known Suzuki polymerisation. In overview, this method involves polymerising monomers having at least two reactive groups selected from boronic acid, boronic acid ester (preferably C1-C6)); borane (preferably C1-C6) with dihalide functional monomers or monomers having one reactive boronic acid, boronic acid ester (preferably C1-C6)), borane (preferably C1-C6) group and one reactive halide functional group with each other in a reaction mixture in the presence of a catalyst. This method is described in US 5,777,070 and a derivative thereof is described in WO/53656. The contents of these documents are incorporated herein by reference.

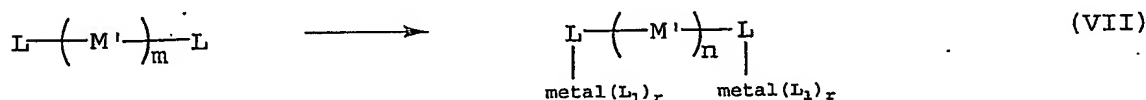
In this regard, a first embodiment of a second aspect of the present invention provides a process for preparing a material as defined in the first aspect of the present invention which comprises:

(a) reacting monomers to form a polymer or oligomer wherein each monomer has at least two reactive groups selected from the group consisting of a halide group, a boronic acid group, a boronic ester group and a borane group and each monomer comprises an aryl or heteroaryl group; and

(b) terminating the polymer or oligomer formed in step (a) using an end-capping reagent, said end-capping reagent comprising one reactive group selected from the group consisting of a halide group, a boronic acid group, a boronic ester group and a borane group and either (i) containing an organometallic as defined above or (ii) being capable of forming a complex with a metal when treated with a metal complexing reagent; and

(a) where the end-capping reagent is as defined in (ii), treating the terminated polymer or oligomer with a metal-complexing reagent.

The treatment in step (c) of the polymer or oligomer formed in step (b)(ii) above may be considered to be as shown in scheme VII:



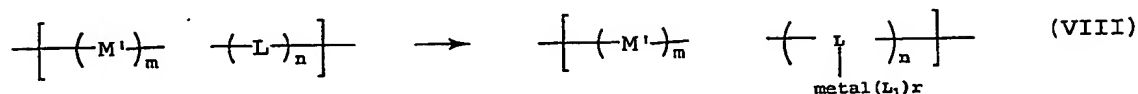
where L, M', m, L' and r are as defined above.

In a second embodiment of the second aspect of the present invention, there is provided a process for preparing a material as defined in the first aspect of the present invention which includes reacting at least one first monomer with a plurality of second monomers which are different to the first monomer to form a polymer or oligomer;

wherein each monomer comprises an aryl or heteroaryl group and has at least two reactive groups selected from the group consisting of a halide group, a boronic acid group, a boronic ester group and a borane group; and wherein the first monomer either (i) contains an organometallic as defined above or (ii) is capable of forming a complex with a metal when treated with a metal complexing reagent; and

(c) where the first monomer is as defined in (ii), treating the polymer or oligomer with a metal-complexing reagent.

The treatment in step (c) of the polymer or oligomer formed in step (b)(ii) may be considered to be as shown in scheme VIII:



where L, M', m, n, L, L₁ and r are as defined above.

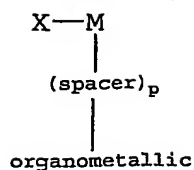
It will be clear from the above methods that, broadly speaking, the organometallic ultimately can be included in the present material as part of an end-capping group or as part of a monomeric residue. Within these two classes of method, the organometallic may be present in the end-capping reagent or monomer when first added to the reaction mixture. Alternatively, it may be introduced in a subsequent step after the terminated polymer or oligomer has been formed. The appropriate method may be selected according to the desirable nature, location and proportion of the polymer or oligomer and the organometallic in the material.

The methods according to the second aspect of the present invention enable manufacture of polymers having general formulas I, II and III.

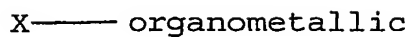
In some embodiments of the second aspect of the present invention, it has been found that ligands sometimes inadvertently are removed from the metal during the reaction. Accordingly, it may be preferable to include a further step in these methods of restoring any removed ligands to the metal.

Surprisingly, the metal itself is not detached from L during the Suzuki reaction.

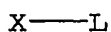
In the first embodiment of the second aspect of the present invention, it is preferred that the end capping reagent has a formula as shown in general formula X, XI or XII.



(X)



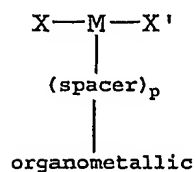
(XI)



(XII)

where X = reactive group as defined above, and L is a ligand capable of forming a complex with a metal when treated with a metal-complexing reagent.

In the second embodiment of the second aspect of the present invention, it is preferred that the first monomer has a general formula as shown in formula IV, V or VI



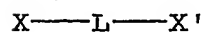
(IV)

(i) $\text{X}=\text{X}'$ (ii) $\text{X} \neq \text{X}'$



(V)

(i) $\text{X}=\text{X}'$ (ii) $\text{X} \neq \text{X}'$



(VI)

(i) $\text{X}=\text{X}'$ (ii) $\text{X} \neq \text{X}'$

where X and X' each is a reactive groups independently selected from the group consisting of a halide group, a boronic acid group, a boronic ester group and a borane group and L is a ligand capable of forming a complex with a metal when treated with a metal-complexing reagent; p o; and M is a group comprising an aryl or heteroaryl group.

Preferably, M is a group selected from the group consisting of 2,7-linked 9,9 disubstituted fluorene, , a p-linked dialkyl phenylene, a p-linked disubstituted phenylene, a phenylene vinylene, a 2,5-linked benzothiadiazole, a 2,5-linked substituted benzothiadiazole, a 2,5-linked disubstituted benzothiadiazole, a 2,5-linked substituted or unsubstituted thiophene or a triarylamine.

Preferably, the first monomer has only two reactive groups to achieve a linear polymer. However, some cross-linking may be achieved by having more than two reactive groups.

In order to control further the structure of the present material, the nature of the reactive groups can be selected so as to control the ordering of the monomers. In one embodiment, each monomer independently can be selected to contain either (a) two reactive halide groups; or (b) two reactive boron derivate groups independently selected from a boronic acid group, a boronic ester group and a borane group. In other embodiment, each monomer can be selected to contain one reactive boron derivate group independently selected from a boronic acid group, a boronic ester group and a borane group and one reactive halide group.

As mentioned above, the present materials will be useful in OLED's. As such, a third aspect of the present invention provides an optical device for a component therefor which comprises a substrate and a material according to the first aspect of the present invention supported on the substrate.

Preferably, the optical device comprises an electroluminescent device. A preferred structure for an electroluminescent device comprises a first charge carrier injecting layer for injecting positive charge carriers, a second charge carrier injecting layer for injecting negative charge carriers, a light-emissive layer located between the charge

carrier injecting layers for generating light and comprising a material according to the first aspect of the present invention.

According to a fourth aspect of the present invention, an electrically pumped organic laser comprising an emissive material according to the first aspect of the present invention also is provided.

The present invention now will be described in more detail with reference to the accompanying figures in which:

Figure 1 shows absorption and PL-spectra of $\text{Ir(ppy-X)}_2(\text{acac})$ complexes;

Figure 2 shows EL-spectra of $\text{Ir(ppy-X)}_2(\text{acac})$;

Figure 3 shows I-L-V curves of $\text{Ir(ppy-X)}_2(\text{acac})$;

Figure 4 shows PL-spectra of blends of complex N in PF8. Insert shows triplet signal of blends of 5.0 mol% of complex L and N in PF8.

Figure 5 shows EL-spectra of $\text{Ir(ppy-X)}_2(\text{acac})$ in polyfluorene;

Figure 6 shows PL-spectra of $\text{Ir(ppy-X)}_2(\text{acac})$ incorporated polyfluorenes;

Figure 7 shows EL-spectra of polyfluorene incorporated Iridium-complexes;

Figure 8 shows EL-spectra of S_2 at increasing driving voltage, going from 8.5V (bottom) up in steps of 0.5V to 12.5V (top);

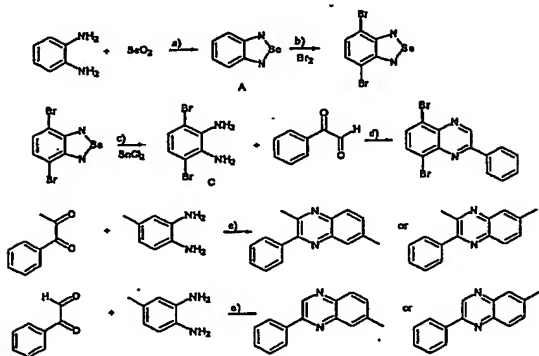
Figure 9 shows scheme 1 as referred to below;

Figure 10 shows scheme 10 as referred to below;

Figure 11 shows scheme 3 as referred to below;

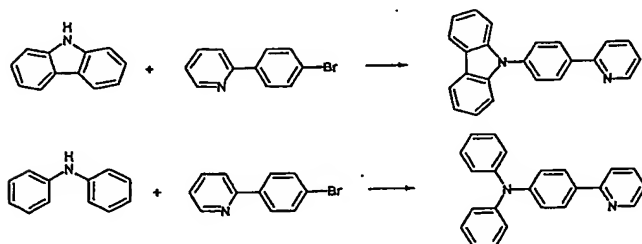
Figure 12 shows scheme 4 as referred to below.

Examples of electron transport materials as possible ligands (L) in the organometallic are shown below:



The above synthetic routes are assembled from synthetic pathways described in literature a) 10 min at 100°C in H₂O/EtOH 1:1. b) 1 h at 20°C in presence of Ag₂SO₄ in conc. H₂SO₄. c) 2h at 70°C in conc. HCl. d) 3h at 100°C in H₂O/EtOH. e) 3h at 100°C in H₂O/EtOH.

Examples of hole transport materials as possible ligands (L) in the organometallic are shown below:

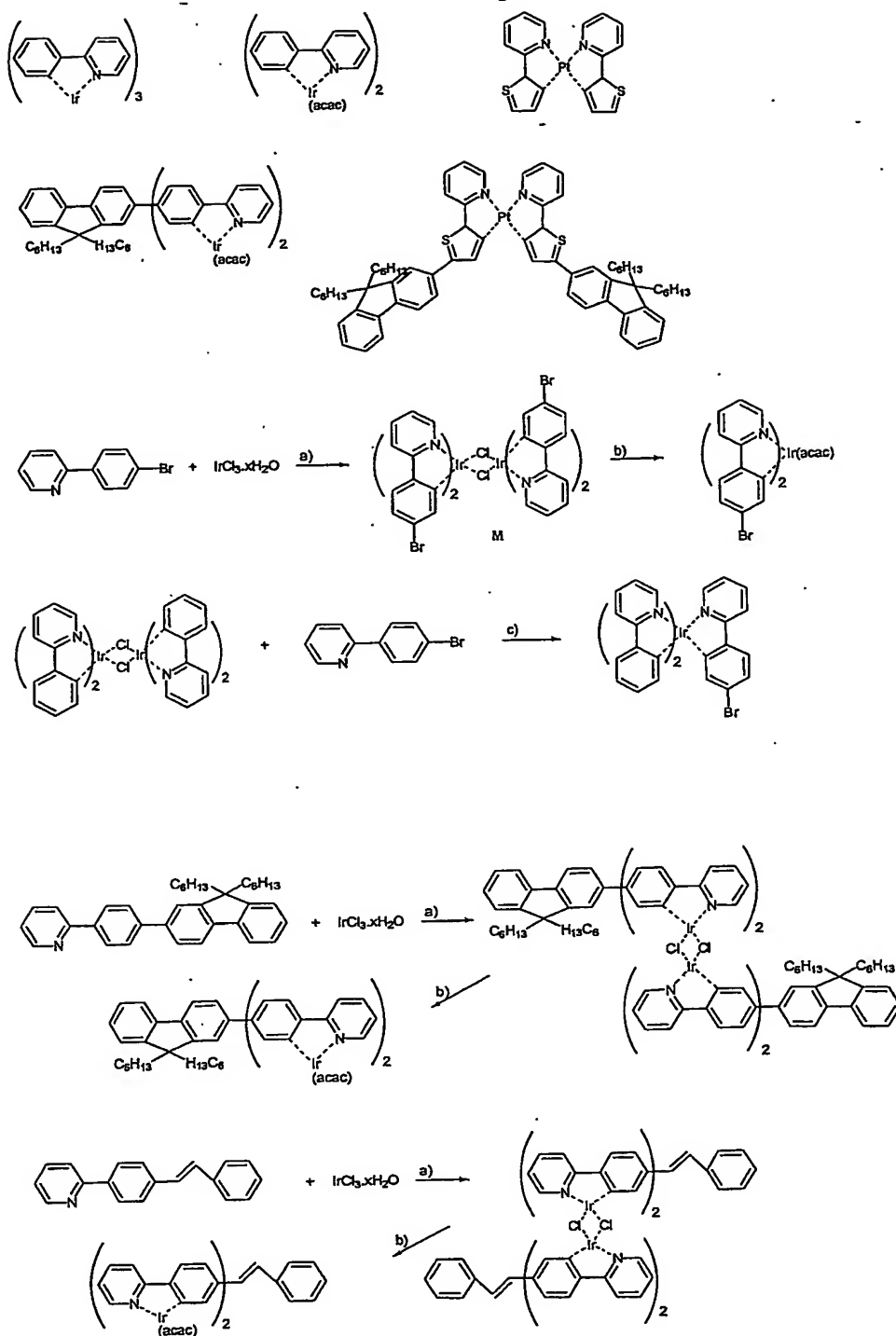


The above reactions were performed in toluene at 90°C for 18 h in the presence of Pd(acetate)/P(tBu)₃ and KOtBu. These ligands were designed with the idea of combining the hole transport ability of the diarylamine units with the ligating possibilities of the phenylpyridyl unit. The triarylaminines were synthesised by a palladium coupling in the presence of a base.

Examples of semi-conducting materials (such as fluorenes or phenylene-vinylene) as part of the ligand system (L) in the organometallic are shown below:.



Some complexes of these monomers are shown below:



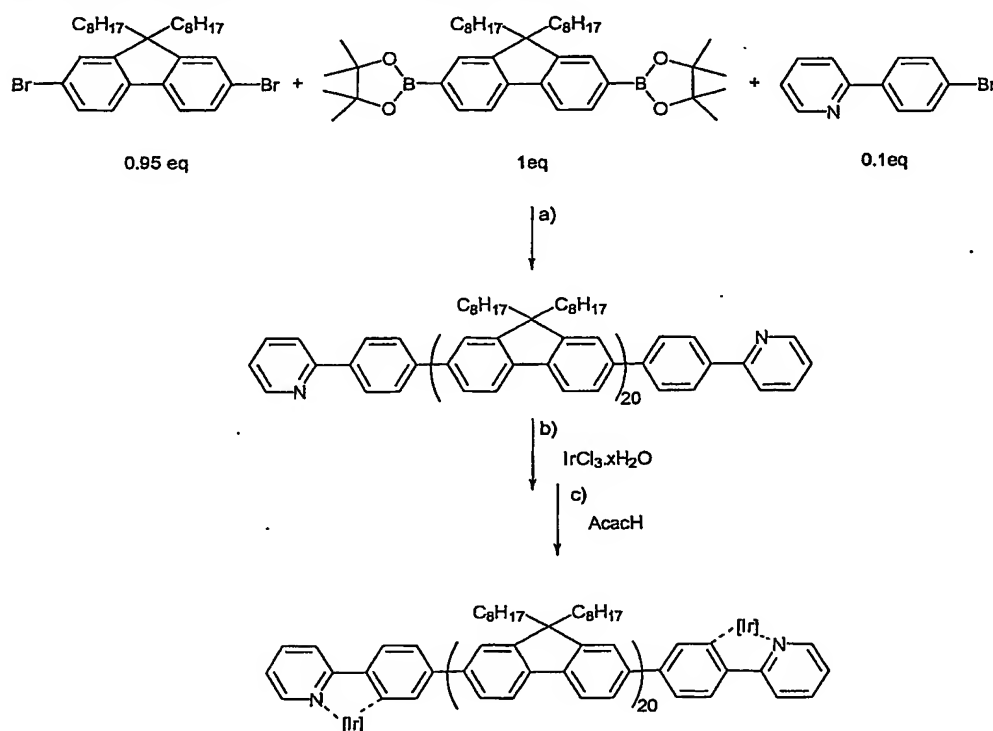
a) Heat reaction mixture to 110-130°C in 2-ethoxyethanol/ H_2O 10:3 for 3-8 h.

b) Heat reaction mixture to 110-130°C in 2-ethoxyethanol in the presence of acetylacetone and Na_2CO_3 .

c) Heat the neat reaction mixture (no solvents) up to 100°C in the presence of AgCF_3SO_3 .

Iridium incorporated polyfluorene was attempted via three different routes.

(1) In the first case the phenylpyridine ligand was incorporated in polyfluorene prior to complexation as described below:

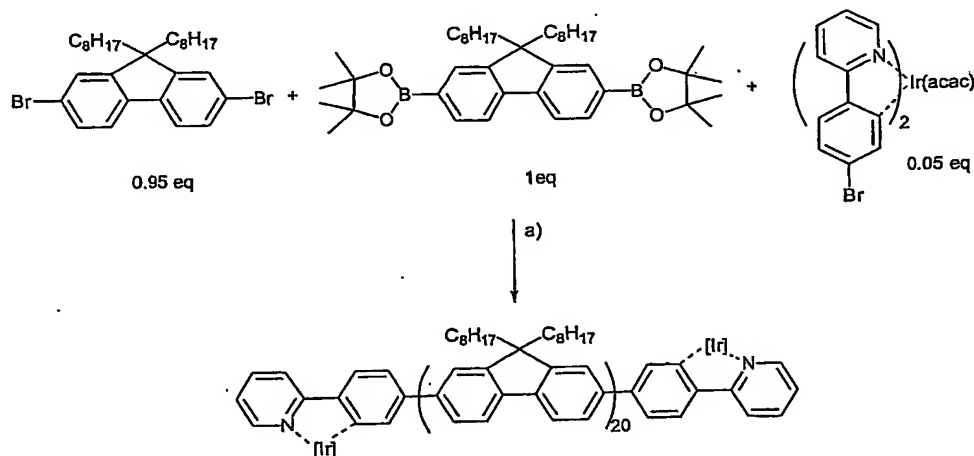


To a solution of 500 mg (0.78 mmol) of 2,7-di(4,4,5,5-tetramethyl-1,3,2-dioxaboronate)-9,9-dioctylfluorene, 405.5 mg (0.740 mmol) of 2,7-dibromo-9,9-dioctylfluorene and 17.1 mg (0.078 mmol) 2-(6-pyridyl)phenyl in 3 ml toluene was added: 4 mg (0.004 mmol) $\text{Pd}(\text{PPh}_3)_4$ and 2.5 ml of (3.1 mmol) Et_4NOH . The resulting mixture was stirred for 24 h at 100°C. After 3h, 5 ml toluene was added to dissolve the precipitated oligomers or to rise the viscosity of the mixture. After the mixture was cooled down to 20°C the mixture was slowly transferred in a large excess of MeOH, allowing the polymer to precipitate as a white (slightly greenish) fibre-like solid. The solids were collected on a filter, redissolved in 50 ml of toluene and subsequently filtered on a short column of silica. The filtrate was transferred into MeOH resulting in a reprecipitation of the polymer as off-white fibres. The solvents were removed under reduced pressure. ^1H NMR (250 MHz, CDCl_3): δ = 8.72 (br d,

2H*12%; CH-N), 8.13 (br d, 2H*12%, CH-CH-CH-N), 7.5-7.9 (br, 12H, ArH), 2.0-2.3 (br, 4H, $\text{CH}_2\text{-C}$), 1.0-1.3 (br, 12H, CH_2), 0.7-0.9 (br, 10H, CH_3+CH_2). GPC: Mp 48586, Mn 18618, Mw 67120, polydispersity 3.6.

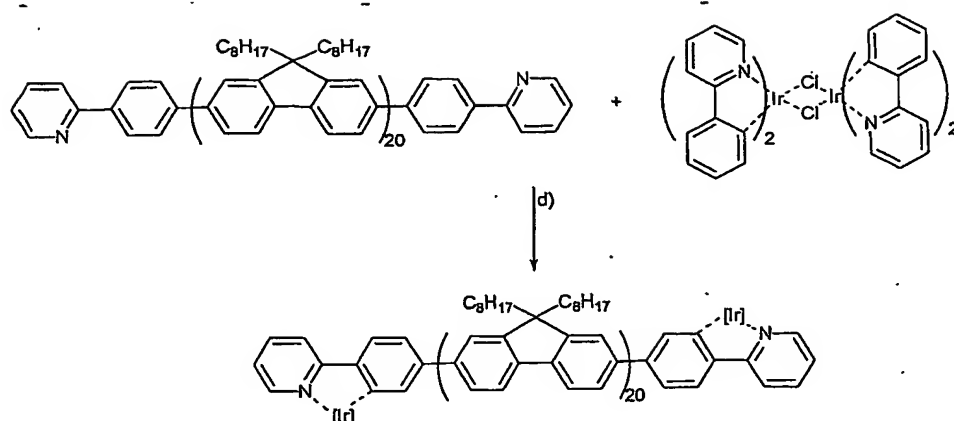
Next the iridium was introduced using the following method: A mixture of 200 mg (containing 0.159 mmol 2-(6-p-pyridyl-phenyl) and 283 mg (containing 0.80 mmol of iridium) was transferred in a solvent mixture of 5 ml of toluene, 3 ml of H_2O and 3 ml of 2-ethoxyethanol. Upon heating the resulting clear dark purple mixture to 90°C the polymer product starts to precipitate after a few hours. After 24 h the water phase was only slightly coloured and the reaction mixture was allowed to cool down to 20°C . Upon addition of 10 ml of H_2O to the mixture the polymer precipitated further as yellow fibres. The complete mixture was transferred in a large excess of MeOH and the precipitated polymer was subsequently collected on a filter, washed with water, methanol and acetone and dried under reduced pressure. Yield: 161 mg.

(2)



In the second approach 0.05 eq of bromide functionalised iridium compound was copolymerised in the Suzuki polymersation of polyfluorene to yield. The method was as follows: A polymer was synthesised starting with 500 mg (0.779 mmol) 2,7-di(4,4,5,5,-tetramethyl-1,3,2,-dioxaboronate)-9,9-dioctylfluorene, 405.5 mg (0.740 mmol) of 2,7-dibromo-9,9-dioctylfluorene and 29.5 mg (0.0389 mmol) (Ir(2(4-bromophenyl)pyridine)2acac).

(3)



In the third route phenylpyridine equipped polyfluorene was reacted with $[\text{Ir}(\text{PhPy})_2\mu\text{Cl}]_2$ in the presence of a silver salt to yield. The method was as follows: A suspension of 180 mg (eq. of 0.143 mmol 2-phenylpyridine) of polyfluorene, 77 mg (0.072 mmol) of $[\text{Ir}(\text{2-phenylpyridine})_2\mu\text{-Cl}]_2$ and 37 mg (0.143 mmol) of AgCF_3SO_3 in 5 ml toluene was heated at 100°C for 18 h. The resulting mixture was directly filtered over a short silica column, eluting with toluene. An off-white polymer was obtained upon removing the solvent from this fraction. The column was further eluted using toluene:MeOH 90:10. Hence a dark green polymer was obtained upon removing the solvents from this fraction under reduced pressure.

EXPERIMENTAL SECTION

To demonstrate the present invention, a study has been carried out on the attachment of subsequent fluorenyl segments to the 4-position of the ligand of a bis-cyclometalated iridium phosphor. The energy levels of the novel phosphors are shown to be complementary to polyfluorene after having attached a monofluorenyl substituent. It is shown furthermore that the stepwise addition of more fluorenyl segments to the phosphor gives rise to a gradual improvement of its performance as the emitting layer in an OLED device structure. The polyaddition to the phosphor was further extended towards the development of a polyfluorenyl-phosphor hybrid system.

Experimentals

General information

All reactions and manipulations were routinely performed under an argon or nitrogen atmosphere using standard Schlenk techniques or a glovebox. Acetonitrile and

dichloromethane were distilled from CaH_2 and toluene and THF were distilled from Na prior to use. Water, glycerol, 2-ethoxyethanol and a 20% Et_4NOH solution in water were degassed for several h before use. All other reagents and chemicals were reagent grade and were used as received from commercial suppliers. Column chromatography was performed using silica gel [Merck 9385 Kieselgel 60 (230-400 ASTM)]. TLC was performed on 0.25 mm thick plates precoated with Merck Kieselgel 60 F₂₅₄ silica gel. ^1H NMR spectra were recorded on Bruker DPX-250 (250 MHz), Bruker DRX-400 (400 MHz) and Bruker DRX-500 (500 MHz) instruments using CDCl_3 or C_6D_6 as an internal deuterium lock. Chemical shifts are in ppm relative to TMS and are indicated as: s(singlet), d(doublet), tr(triplet), q(quartet), br(broad), m(multiplet), dd(double doublet), dtr(double triplet) etc. ^{13}C NMR spectra were recorded on Bruker DPX-250 (62.5 MHz) and Bruker DRX-400 (100 MHz) and a Bruker DRX-500 (121 MHz) instruments using an internal deuterium lock and proton decoupling. The multiplicity of the signal was determined by APT (Attached Proton Test) experiments. Mass spectra were recorded by the Mass Spectrometry Services in the University of Swansea. Electron Impact (IE) and Chemical Ionisation (CI) low resolution spectra were carried out on a VG model 12-253 under ACE conditions. Accurate mass measurements for EI and CI were performed on a +VG ZAB-E instrument. Melting points were determined using a Büchi 510 melting point apparatus, and are uncorrected.

Synthesis of A (2(4-bromophenyl)pyridine)^{1,2}

At 0°C, a solution of 16.84 g (244 mmol) NaNO_2 in 30 ml H_2O was slowly added to a suspension of 20 g (116.4 mmol) p-bromoaniline in 40 ml concentrated HCl. The resulting mixture was stirred for an h at below 5°C and was subsequently carefully poured in 500 ml pyridine. The resulting clear brown reaction mixture was stirred at 40°C for 4 h. 200 g of Na_2CO_3 was added to the mixture and the slurry was stirred for 18 h at 40°C. The organic layer was dissolved in dichloromethane, extracted with water and dried with MgSO_4 . The crude product was purified by flash column chromatography (toluene:MeOH 95:5). Yield: 8.4 g (33%), considerable yields of 7-(p-bromophenyl)pyridine were obtained as the major side product. ^1H NMR (500 MHz, CDCl_3): δ = 8.68 (d, 3J = 4.3 Hz, 1H; CH-N), 7.88 (d, 3J = 8.4 Hz, 2H; CH-CH-Br), 7.75 (d, 3J = 7.5 Hz, 1H; CH-CH-CH-N), 7.70 (d, 3J = 7.8 Hz, 1H; CH-CH-CH-

CH-N), 7.60 (d, $^3J = 8.4$ Hz, 2H, CH-Br), 7.24 (br, 1H, CH-CH-N). $^{13}\text{C}\{^1\text{H}\}$ NMR (500 MHz, CDCl_3): $\delta = 156.2$ (CN), 149.8 (CH-N), 138.3 (C-CN), 136.8 (CH-CH-CH-N), 131.9, 128.5 (Ar-CH), 123.4 (CBr), 122.4 (CH-CN), 120.2 (CH-CH-N). Exact Mass (IE) m/z : 233.9920 $[\text{M}+\text{H}]$ (calcd. 233.9918 (^{79}Br)). $\text{C}_{11}\text{H}_8\text{BrN}$: calcd. C 56.44, H 3.44, N 5.98; found C 56.24, H 3.35, N 5.90.

Synthesis of B 2-(4,4,5,5 tetramethyl-1,3,2-dioxaborolane)-7-trimethylsilyl-9,9-dioctylfluorene

At -78°C , 7.3 mL of a 1.7 M (12.5 mmol) tBuLi solution in hexanes was slowly added to a solution of 2.7 g (4.99 mmol) 2-bromo-7-trimethylsilyl-9,9-dioctylfluorene in THF. After 1 h, 1.2 g (6.48 mmol) of 2-isopropoxy-4,4,5,5 tetramethyl-1,3,2-dioxaborolane was added to the green solution at -78°C , after which the reaction mixture was allowed to warm up to room temperature. The resulting red solution turned colourless upon stirring for 18 h at room temperature. The product was extracted using dichloromethane/water and the crude product was purified by flash column chromatography (silica, 3% CDM in hexane). Yield 1.2 g (41%) product was obtained as colourless oil. ^1H NMR (500 MHz, CDCl_3): $\delta = 7.80$ (d, $^3J = 7.5$ Hz, 1H; ArH), 7.75 (s, 1H; ArH), 7.70 (d, $^3J = 7.5$ Hz, 2H; ArH), 7.48 (d, $^3J = 7.5$ Hz, 1H; ArH), 7.46 (s, 1H, ArH), 1.97 (tr, 4H, $^3J = 8.3$ Hz, $\text{CH}_2\text{-CH}_2\text{-C}$), 1.39 (s, 12H, $\text{CH}_3\text{-C(O)}$), 1.05-1.15 (mp, 20H, CH_2), 0.81 (t, $^3J = 7.1$ Hz, 6H; $\text{CH}_3\text{-CH}_2$), 0.61 (mp, 4H; CH_2), 0.31 (s, 9H, $\text{CH}_3\text{-Si}$). $^{13}\text{C}\{^1\text{H}\}$ NMR (500 MHz, CDCl_3): $\delta = 150.4$ (C-C-CH_2), 150.1 (C-C-CH_2), 144.1, 141.5, 139.6 (C), 133.6, 131.7, 128.9, 127.6, 119.3, 119.0 (CH), 83.7 (CO), 55.0 (C-CH_2), 40.0 (C-CH_2), 31.7, 30.0, 29.1, 29.0 (CH_2), 24.9 ($\text{CH}_3\text{-CO}$), 23.6 ($\text{C-CH}_2\text{-CH}_2$), 22.6 ($\text{CH}_3\text{-CH}_2$), 14.0 ($\text{CH}_3\text{-CH}_2$), 0.9 ($\text{CH}_3\text{-Si}$). Exact Mass (IE) m/z : 588.4534 (calcd 588.4534), fractions: 475.4 (m; $[\text{M}-(\text{C}_8\text{H}_{17})]^+$), 363.3 (w; $[\text{M}-(\text{C}_8\text{H}_{17})_2]^+$), 333.3 (m; $[\text{M}-(\text{C}_8\text{H}_{17}+\text{CH}_3+\text{C}_6\text{H}_{12}\text{BO}_2)]^+$). $\text{C}_{38}\text{H}_{71}\text{BO}_2\text{Si}$: calcd. C 77.52, H 10.44; found C 77.35, H 10.33.

Synthesis of C 2-(4,4,5,5 tetramethyl-1,3,2-dioxaborolane)-7-bromo-9,9-dioctylfluorene

At -78°C , 4.8 mL of a 1.6M (7.66 mmol) nBuLi solution in hexanes was slowly added to a solution of 4 g (7.30 mmol) of 2,7-dibromo-9,9-dioctylfluorene in 50 mL of diethylether. The resulting white milky solution was allowed to cool to 0°C until a

clear-yellow solution was formed that was cooled back to -78°C again. 1.49 g (8.03 mmol) of 2-isopropoxy-4,4,5,5-tetramethyl-1,3,2-dioxaborolane was added and the resulting mixture turned white turbid after 1 h. The mixture was subsequently allowed to cool to room temperature and stirred for 18 h. After the mixture was quenched with water, the organic layer was dried on MgSO_4 and the solvent was removed under reduced pressure. The crude product was purified using flash column chromatography (silica, dichloromethane, Rf: 0.5). Yield: 2.2 g (51%). ^1H NMR (500 MHz, CDCl_3): δ = 7.80 (d, 3J = 7.6 Hz, 1H; ArH), 7.72 (s, 1H; ArH), 7.66 (d, 3J = 7.5 Hz, 1H; ArH), 7.57 (d, 3J = 7.5 Hz, 1H; ArH), 7.47 (s, 1H; ArH), 7.45 (d, 3J = 7.5 Hz, 1H; ArH), 1.97 (mp, 2H, $\text{CH}_2\text{-CH}_2\text{-C}$), 1.91 (mp, 2H, $\text{CH}_2\text{-CH}_2\text{-C}$), 1.39 (s, 12H, $\text{CH}_3\text{-C(O)}$), 1.02-1.18 (mp, 20H, CH_2), 0.82 (t, 3J = 7.1 Hz, 6H; $\text{CH}_3\text{-CH}_2$), 0.56 (mp, 4H; CH_2). $^{13}\text{C}\{^1\text{H}\}$ NMR (500 MHz, CDCl_3): δ 153.3, 149.2 (C-C-CH_2), 142.7, 139.7 (ArC), 127.9 (very weak) (C-B), 133.6, 129.6, 128.6, 126.0, 121.2, 118.8 (CH), 83.5 (CO), 55.2 (C-CH_2), 39.9 (C-CH_2), 31.5, 29.6, 28.91, 28.88 (CH_2), 24.6 (C-CH_3), 23.4 ($\text{C-CH}_2\text{-CH}_2$), 22.3 ($\text{CH}_2\text{-CH}_3$), 13.8 ($\text{CH}_2\text{-CH}_3$). $\text{C}_{35}\text{H}_{52}\text{BBrO}_2$: calcd. C 70.59, H 8.80; found C 70.25, H 8.47.

Synthesis of **D** (2-(6-pyridyl-phenyl)-9,9-dihexylfluorene)

A mixture of 500 mg (1.12 mmol) of 2-(4,4,5,5-tetramethyl-1,3,2-dioxaboronate)-9,9-dihexylfluorene, 261 mg (1.12 mmol) of 2-(4-bromophenyl)pyridine, 0.5 mg (0.01 mmol) $\text{Pd(PPh}_3)_4$, 2 ml of a 20% Et_4NOH in water in 4 ml toluene was stirred at 90°C for 18 h. The clear brown-orange mixture was taken in dichloromethane, washed with water and finally dried with MgSO_4 . The crude product was purified by flash column chromatography (DCM). Yield 255 mg (47%). ^1H NMR (250 MHz, CDCl_3): δ = 8.73 (d, 3J = 4.9 Hz, 1H; CH-N), 8.12 (d, 3J = 8.2 Hz, 2H; $\text{CH-C-C(CH}_3\text{)-N}$), 7.80-7.78 (mp, 4H, ArH), 7.73 (d, 3J = 7.8 Hz, 1H; CH-CH-CH-N), 7.64 (d, 3J = 7.9, 1H; CH-CH-N), 7.62 (br, 1H; CH-CH-N), 7.36-7.33 (mp, 3J = 1.0 Hz and 7.5 Hz, 3H; ArH), 7.25 (mp, 2H, ArH), 2.00-2.04 (dd, 3J = 5.0 Hz and 5.2 Hz, 4H; $\text{CH}_2\text{-C}$), 1.05-1.15 (mp, 12H, CH_2), 0.78 (t, 3J = 6.7 Hz, 6H; CH_3), 0.68 (mp, 4H; CH_2). $^{13}\text{C}\{^1\text{H}\}$ NMR (500 MHz, CDCl_3): δ = 156.8 (CN), 151.2, 150.8 (C-C-CH_2), 149.5 (CH-N), 142.0, 140.5, 139.1, 137.8 (ArC), 136.5 (CH-CH-CH-N), 127.2, 127.0, 126.8, 126.5, 126.0, 122.7, 121.8, 121.1, 120.2, 119.7, 119.5 (CH), 54.9 (C-CH_2), 40.2 (C-CH_2), 31.2, 29.5 (CH_2), 23.5 ($\text{C-CH}_2\text{-CH}_2$), 22.3 ($\text{CH}_3\text{-CH}_2$), 13.7 (CH_3). Exact Mass (IE): 488.3317

[M+H] (calcd. 488.3317); fractions m/z: 402.3 (s, [M-(C₆H₁₃)]⁺), 332.2 (s, [M-(C₆H₁₃)-(C₅H₁₁)]⁺), 318.2 (s, [M-(C₆H₁₃)₂]⁺). C₃₆H₄₁N: calcd. C 88.65, H 8.47, N 2.87; found C 88.37, H 8.40, N 3.11.

Synthesis of **E** Suzuki polymerisation of 9,9-di-n-octylfluorene containing 2-(6-pyridyl-phenyl) endgroup on one side

To a solution of 500 mg (0.56 mmol) of 2-(4,4,5,5-tetramethyl-1,3,2-dioxaboronate)-7-bromo-9,9-dioctylfluorene and 13.1 mg (0.056 mmol) 2-(6-pyridyl-phenyl) in 5 ml toluene was added: 4 mg (0.004 mmol) Pd(PPh₃)₄ and 2.5 ml of a 20% (3.1 mmol) Et₄NOH solution in water. The resulting mixture was stirred for 24 h at 110°C. After the mixture was cooled down to 20°C the mixture was slowly transferred in a large excess of MeOH, allowing the polymer to precipitate as a yellow-white solid. The solids were collected on a filter, redissolved in 50 ml of toluene and subsequently filtered on a short column of silica. The filtrate was transferred into MeOH resulting in a reprecipitation of the polymer as off-white fibres. The solvents were removed under reduced pressure. ¹H NMR (250 MHz, CDCl₃): δ = 8.75 (br d, 2H*12%; CH-N), 8.13 (br d, 2H*12%, CH-CH-CH-N), 7.5-7.9 (br, 6H, ArH), 2.0-2.3 (br, 4H, CH₂-C), 1.0-1.3 (br, 20H, CH₂), 0.7-0.9 (br, 10H, CH₃+CH₂). Mw.(NMR): 5000 (12-mer).

Synthesis of **F** 2-(6-p-pyridyl-phenyl)-7-trimethylsilyl-9,9-dioctylfluorene

To a solution of 243 mg (1.021 mmol) of (2(4-bromophenyl)pyridine), 600 mg (1.021 mmol) of 2-(4,4,5,5 tetramethyl-1,3,2-dioxaborolane)-7-trimethylsilyl-9,9-dioctylfluorene in 10 mL of toluene was added: 4 mg (0.004 mmol) Pd(PPh₃)₄ and 2.5 ml of a 20 % (3.1 mmol) Et₄NOH solution in water. This two-phase reaction mixture was stirred for 48 h at 110°C after which it was extracted using dichloromethane/water. The organic layer was dried using MgSO₄ and the solvents were removed under reduced pressure. The crude product was purified using flash chromatography (dichloromethane, R_f: 0.47). 550 mg (88%) product was obtained as an off-white solid. ¹H NMR (250 MHz, CDCl₃): δ = 8.73 (d, ³J = 4.9 Hz, 1H; CH-N), 8.12 (d, ³J = 8.3 Hz, 2H; CH-C-C(CH₃)-N), 7.80-7.78 (mp, 4H, ArH), 7.73 (d, ³J = 4.5 Hz, 2H, CH-CH-CH-N), 7.69 (d, ³J = 4.6 Hz, 2H; CH-CH-N), 7.63 (d, ³J = 8.0 Hz, 2H; CH-CH-N), 7.25 (s, 1H; ArH), 2.01 (tr, ³J = 8.2 Hz, 4H; CH₂-C), 1.05-1.15 (mp, 20H, CH₂), 0.78 (t, ³J = 6.6 Hz, 6H; CH₃), 0.70 (mp, 4H; CH₂), 0.32 (s, 9H; CH₃-Si).

$^{13}\text{C}\{^1\text{H}\}$ NMR (500 MHz, CDCl_3): δ = 156.8 (CN), 151.5, 149.9 ($\text{C}-\text{C}-\text{CH}_2$), 149.5 ($\text{CH}-\text{N}$), 142.0, 141.1, 140.4, 139.2, 138.9, 137.8 (ArC), 136.5 ($\text{CH}-\text{CH}-\text{CH}-\text{N}$), 131.6, 127.4, 127.2, 127.0, 125.7, 121.8, 121.2, 120.1, 119.8, 118.8 (CH), 54.9 ($\text{C}-\text{CH}_2$), 39.9 ($\text{C}-\text{CH}_2$), 31.5, 28.9, 28.8 (CH_2), 29.7 ($\text{C}-\text{CH}_2-\text{CH}_2-\text{CH}_2$), 23.5 ($\text{C}-\text{CH}_2-\text{CH}_2$), 22.3 (CH_3-CH_2), 13.8 (CH_2-CH_3), -1.1 ($\text{Si}-\text{CH}_3$). Exact Mass (IE): $[\text{M}+\text{H}]^+$: 616.4340 (calcd. 616.4338). $\text{C}_{43}\text{H}_{57}\text{NSi}$: calcd. C 83.84, H 9.33, N 2.27; found C 83.99, H 9.41, N 2.33.

Synthesis of **G** 2-(6-p-pyridyl-phenyl)-7-iodo-9,9-dioctylfluorene

1.5 ml of an 1.0 M ICl solution in MeOH was slowly added to a solution of 500 mg (0.813 mmol) of 2-(6-p-pyridyl-phenyl)-7-trimethylsilyl-9,9-dioctylfluorene in 5 mL of dichloromethane. The resulting darkred solution was stirred for 2 h at roomtemperature after which the reactionmixture was quenched with a large excess of a $\text{Na}_2\text{S}_2\text{O}_3$ in water. The mixture was extracted with water, and the organic fraction was dried using MgSO_4 and the solvents were removed under reduced pressure. The product was purified using flash column chromatography (dichloromethane: Rf: 0.36). 450 mg (85%) product was obtained as a highly blue fluorescent, colourless solid material. ^1H NMR (250 MHz, CDCl_3): δ = 8.73 (d, 3J = 4.7 Hz, 1H; CH-N), 8.12 (d, 3J = 8.5 Hz, 2H; $\text{CH}-\text{C}-\text{C}(\text{CH})-\text{N}$), 7.80-7.73 (mp, 4H, ArH), 7.68 (s, 2H, ArH), 7.65 (d, 3J = 5.0 Hz, 2H; $\text{CH}-\text{CH}-\text{N}$), 7.60 (s, 2H; ArH), 7.48 (d, 1H; 3J = 8.4 Hz; ArH), 2.05-1.94 (mp, 4H; CH_2-C), 1.05-1.25 (mp, 20H, CH_2), 0.81 (t, 3J = 6.6 Hz, 6H; CH_3), 0.70 (br m, 4H, CH_2). $^{13}\text{C}\{^1\text{H}\}$ NMR (500 MHz, CDCl_3): δ = 157.0 (CN), 153.4, 150.9 ($\text{C}-\text{C}-\text{CH}_2$), 149.8 (CH-N), 142.0, 140.4, 140.0, 139.6, 138.2 (ArC), 136.8 ($\text{CH}-\text{CH}-\text{CH}-\text{N}$), 135.9, 132.1 ($\text{C}-\text{Cl}$), 127.5, 127.3, 126.1, 122.1, 121.5, 121.3, 120.4, 120.1 (CH), 92.6 (Cl), 55.5 ($\text{C}-\text{CH}_2$), 40.2 ($\text{C}-\text{CH}_2$), 31.7, 29.9, 29.2, 29.1 (CH_2), 23.7 ($\text{C}-\text{CH}_2-\text{CH}_2$), 22.6 (CH_3-CH_2), 14.1 (CH_3). Exact Mass (IE) m/z: 669.2825 $[\text{M}]^+$ (calcd. 669.2832) fractions: 543.6 (m; $[\text{M}-\text{I}]^+$), 318.4 (m; $[\text{M}-(\text{C}_8\text{H}_{17})-\text{I}]^+$). $\text{C}_{40}\text{H}_{48}\text{IN}$: calcd. C 71.74, H 7.22, N 2.09; found C 72.34, H 7.20, N 2.12.

Synthesis of **H** 2-(6-p-pyridyl-phenyl)-7-(9,9-dihexylfluorenyl)-9,9-dioctylfluorene

To a solution of 400 mg (0.60 mmol) of 2-(6-p-pyridyl-phenyl)-7-iodo-9,9-dioctylfluorene, 362 mg (0.60 mmol) of 2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolane)-

9,9-dihexylfluorene in 10 mL of toluene was added: 4 mg (0.004 mmol) $\text{Pd}(\text{PPh}_3)_4$ and 2.5 ml of an 20 % (3.1 mmol) Et_4NOH solution in water. This two-phase reactionmixture was stirred for 48 h at 110°C after which it was extracted using dichloromethane/water. The organic layer was dried on MgSO_4 and the solvents were removed under reduced pressure. The crude product was purified using flash chromatography. Yield 450 mg (86%). ^1H NMR (400 MHz, CDCl_3): δ = 8.74 (d, 3J = 4.9 Hz, 1H; CH-N), 8.13 (d, 3J = 4.6 Hz, 2H; CH-C-C(CH)-N), 7.84-7.77 (mp, 6H, ArH), 7.74 (d, 3J = 6.0 Hz, 2H, CH-CH-CH-N), 7.69-7.63 (mp, 6H; ArH), 7.39-7.30 (mp, 3H; ArH), 7.27-7.24 (mp, 1H (obscured by CDCl_3 peak ; ArH), 2.11-2.02 (mp, 8H, C-CH₂), 1.18-1.09 (mp, 32H, CH₂), 0.81-0.74 (mp, 20H, CH₂-CH₃). $^{13}\text{C}\{^1\text{H}\}$ NMR (500 MHz, CDCl_3): δ = 157.1 (CN), 151.8, 151.4, 151.0 (C-C-CH₂), 149.8 (CH-N), 142.2, 140.8, 140.6, 140.42, 140.39, 140.3, 139.9, 139.3, 138.1 (ArC), 136.8 (CH-CH-CH-N), 127.5, 127.3, 126.9, 126.8, 126.1, 126.0, 122.1, 122.0, 121.44, 121.38, 120.4, 120.0, 119.8, 119.7 (CH), 55.3, 55.1 (C-CH₂), 40.4 (C-CH₂), 31.7, 31.5, 30.0, 29.7, 29.18, 29.16 (CH₂), 23.8, 23.7 (C-CH₂-CH₂), 22.6, 22.5 (CH₃-CH₂), 14.03, 13.99 (CH₃). Exact Mass (IE): 875.6367 $[\text{M}]^+$ (calcd 875.6369) fractions m/z:: 762.3 (m; $[\text{M}-(\text{C}_8\text{H}_{17})]^+$, 650.4 (m; $[\text{M}-(\text{C}_8\text{H}_{17})_2]^+$. $\text{C}_{65}\text{H}_{81}\text{N}$: calcd. C 89.09, H 9.32, N 1.60; found: C 88.60, H 9.25, N 1.52.

Synthesis of J (via I) 2-(6-p-pyridyl-phenyl)-7-(2-(7-iodo-9,9-dioctylfluorenyl)-9,9-dioctylfluorene

To a solution of 300 mg (0.448 mmol) of 2-(6-p-pyridyl-phenyl)-7-iodo-9,9-dioctylfluorene, 320 mg (0.544 mmol) of 2-(4,4,5,5 tetramethyl-1,3,2-dioxaborolane)-7-trimethylsilyl-9,9-dioctylfluorene in 10 mL of toluene was added: 4 mg (0.004 mmol) $\text{Pd}(\text{PPh}_3)_4$ and 2.5 ml of a 20 % (3.1 mmol) Et_4NOH solution in water. This two-phase reactionmixture was stirred for 24 h at 110°C after which it was extracted using dichloromethane/water. The organic layer was dried and the solvents were removed under reduced pressure. The crude product was purified using flash chromatography (dichloromethane, Rf: 0.30). 392 mg (88%) of product I was obtained as a colourless solid. This was subsequently dissolved in 5 ml of dichloromethane and 5 mL of 1.0 M solution of ICl in MeOH was added. The

resulting darkred solution was stirred for 2 h at roomtemperature after which the reactionmixture was queched with a large excess of a $\text{Na}_2\text{S}_2\text{O}_3$ solution in water. The reactionmixture was subsequently extracted (dichloromethane/water). The organic phase was dried on MgSO_4 and the solvents were removed under reduced pressure. The crude product was purified using column chromatography (silica, dichloromethane). Yield: 355 mg (85%). ^1H NMR (400 MHz, CDCl_3): δ = 8.74 (d, 3J = 4.6 Hz, 1H; CH-N), 8.13 (d, 3J = 8.4 Hz, 2H; CH-C-C(CH)-N), 7.82-7.78 (mp, 5H, ArH), 7.75 (d, 3J = 7.9 Hz, 2H, CH-CH-CH-N), 7.69-7.64 (mp, 5H; ArH), 7.60 (d, 2H; ArH), 7.48 (d, 1H; CH-CH-N) 7.26-7.24 (mp, 2H (obscured by CDCl_3 peak ; ArH), 2.17-1.94 (mp, 8H, C- CH_2), 1.24-1.09 (mp, 40H, CH_2), 0.90-0.71 (mp, 20H, CH_2 - CH_3). $^{13}\text{C}\{^1\text{H}\}$ NMR (500 MHz, CDCl_3): δ = 157.0 (CN), 153.4, 151.83, 151.77, 150.9 (C-C- CH_2), 149.8 (CH-N), 142.1, 141.1, 140.5, 140.4, 140.3, 140.1, 139.4, 139.3, 138.1 (ArC), 136.8 (CH-CH-CH-N), 135.9, 132.1 (C-Cl), 127.5, 127.3, 126.24, 126.16, 126.1, 122.1, 121.4, 121.3, 120.4, 120.1, 120.0 (CH, 3 CH signals obscured by exact overlap), 92.5 (Cl), 55.5, 55.3 (C- CH_2), 40.4, 40.2 (C- CH_2), 31.77, 31.76, 29.19, 29.17 (CH_2), 29.99, 29.93 (C- CH_2 - CH_2 - CH_2) 23.8, 23.7 (C- CH_2 - CH_2), 22.60, 22.58 (CH_3 - CH_2), 14.08, 14.05 (CH_3). Exact Mass (IE): $[\text{M}+\text{H}]$ (calcd.); fractions m/z: 1058.7 (s, $[\text{M}]^+$), 944.4 (m, $[\text{M}-(\text{C}_8\text{H}_{17})]^+$). $\text{C}_{69}\text{H}_{88}\text{IN}$: calcd. C 78.30, H 8.38, N 1.32; found. C 78.02, H 8.31, N 1.35.

Synthesis of K 2-(6-p-pyridyl-phenyl)-7-(2-(7-[2-(9,9-dihexylfluorenyl)])-9,9-dioctylfluorenyl)-9,9-dioctylfluorene

To a solution of 300 mg (0.287 mmol) of 2-(6-p-pyridyl-phenyl)-7-(2-(7-iodo-9,9-dioctylfluorenyl)-9,9-dioctylfluorene, 225 mg (0.373 mmol) of 2-(4,4,5,5 tetramethyl-1,3,2-dioxaborolane)-7-trimethylsilyl-9,9-dioctylfluorene in 5 mL of toluene was added: 4 mg (0.004 mmol) $\text{Pd}(\text{PPh}_3)_4$ and 2.5 ml of a ?% (3.1 mmol) Et_4NOH solution in water. This two-phase reactionmixture was stirred for 24 h at 110°C after which it was extracted using dichloromethane/water. The organic layer was dried and the solvents were removed under reduced pressure. The crude product was purified using flash chromatography (silica/dichloromethane:hexane 1:1, Rf: 0.24). Yield: 190 mg (52%) of a colourless, highly blue fluorescent solid. ^1H NMR (250 MHz, CDCl_3): δ = 8.75 (d, 3J = 4.6 Hz, 1H; CH-N), 8.14 (d, 3J = 8.4 Hz, 2H; CH-C-C(CH)-

N), 7.86-7.73 (mp, 10H, ArH), 7.74-7.62 (mp, 10H, ArH), 7.40-7.31 (mp, 3H, ArH), 7.28-7.23 (mp, 1H (obscured by CDCl_3), ArH), 2.17-2.02 (mp, 12H, C- CH_2), 1.25-1.01 (mp, 52H, CH_2), 0.90-0.64 (mp, 30H, CH_2 - CH_3).

$^{13}\text{C}\{^1\text{H}\}$ NMR (500 MHz, CDCl_3): δ = 157.1 (CN), 151.8, 151.4, 151.0 (C-C-CH_2), 149.7 (CH-N), 142.2, 140.8, 140.6, 140.5, 140.47, 140.44, 140.39, 140.30, 139.95, 139.90 (ArC), 136.7 (CH-CH-CH-N), 127.4, 127.3, 126.8, 126.1, 126.0, 122.9, 121.44, 121.40, 120.38, 120.0, 119.9, 119.8, 119.7 (CH), 55.33, 55.31, 55.1 (C-CH_2), 40.3 (C- CH_2), 31.7, 31.4, 30.0, 29.7, 29.2 (CH_2), 23.9, 23.8 (C- CH_2 - CH_2), 22.6, 22.5 (CH_3 - CH_2), 14.01, 13.97 (CH_3). Mass (IE) m/z ; 1265 $[\text{M}+\text{H}]^+$. $\text{C}_{94}\text{H}_{121}\text{N}$: calcd. C 89.25, H 9.64, N 1.11; found C 89.25, H 9.71, N 1.21.

Synthesis of $[\text{Ir}(\text{2(4-bromophenyl)pyridine})_2\mu\text{-Cl}]_2^3$

A mixture of 500 mg (1.42 mmol Ir) $\text{IrCl}_3 \cdot x\text{H}_2\text{O}$ and 1.4 g (6.37 mmol) of 2(4-bromophenyl)pyridine in 50 ml 2-ethoxyethanol and 15 ml H_2O was heated to 110°C for 18h. A yellow precipitate started to form after 1h. After the mixture was allowed to cool down to 20°C , the precipitate was collected on a fritted glass filter, washed with $\text{EtOH}:\text{H}_2\text{O}$ 95:5, dissolved in acetone and dried on MgSO_4 . The solvent was removed under reduced pressure. Yield 639 mg (65%). ^1H NMR (250 MHz, CDCl_3): δ = 9.12 (d, 3J = 5.7 Hz, 4H; CH-N), 7.86 (d, 3J = 4.5 Hz, 4H; CH-CH-CBr), 7.80 (dtr, 4J = 1.5 Hz and 3J = 7.2 Hz, 4H; CH- CH-Br), 7.38 (d, 3J = 9.4 Hz, 4H; CH-CH-CH-N), 7.02 (dd, 3J = 1.9 Hz and 9.0 Hz, 4H; CH-C(C)N), 6.84 (dtr, 4J = 1.7 Hz and 3J = 5.6 Hz, 4H; CH-CH-N), 5.94 (d, 4J = 1.9 Hz, 4H; CH-C(C)-Ir). Mass (FAB): m/z 1387 (w, $[\text{M}]^+$), 1352, (w, $[\text{M-Cl}]^+$), 659 (s, $[\text{Ir}(\text{A})_2]^+$). $\text{C}_{44}\text{H}_{32}\text{Br}_4\text{Cl}_2\text{Ir}_2\text{N}_4$: calcd. C 38.08, H 2.03, N 4.04; found C 37.73, H 2.10, N 4.04.

Synthesis of **M** ($[\text{Ir}(\text{2(4-bromophenyl)pyridine})_2\text{acac}]^4$

300 mg (0.216 mmol) of $[\text{Ir}(\text{2(4-bromophenyl)pyridine})_2\mu\text{-Cl}]_2$ was dissolved in 30 ml 2-ethoxyethanol in the presence of 108 mg (1.08 mmol) acacH and 216 mg (2.04 mmol) Na_2CO_3 . The resulting yellow suspension was stirred at 110°C for 8h. After the mixture was allowed to cool down to 20°C 10 ml of H_2O was added. The yellow precipitate was collected on a fritted glass filter, washed with H_2O , hexane and with cold Et_2O , and dried under reduced pressure. Yield: 190 mg (58%). ^1H NMR (250

MHz, CDCl₃): δ = 8.44 (d, 3J = 5.5 Hz, 2H; CH-N), 7.83 (d, 3J = 8.5 Hz, 2H, CH-CH-CH-N), 7.78 (mp, 3J = 5.5 Hz and 4J = 1.5 Hz, 2H; CH-CH-N), 7.41 (d, 3J = 8.3 Hz, 2H; CH-CH-CH-CH-N), 7.19 (dtr, 3J = 6.4 Hz and 4J = 1.5 Hz, 2H, ArH), 7.00 (dd, 3J = 6.3 Hz and 4J = 2 Hz, 2H; ArH), 6.28 (d, 4J = 2 Hz, 2H; CH-C(C)-Ir), 5.23 (s, 1H; CH-C=O), 1.79 (s, 6H; CH₃). ¹³C{¹H} NMR (500 MHz, CDCl₃): ¹³C{¹H} NMR (500 MHz, CDCl₃): δ = 184.8 (CO), 167.5 (CN), 148.2 (CH-N), 143.8 (C-Ir), 137.3, 135.4 (ArC), 125.1, 124.1, 121.9, 118.7 (Ar-CH), 100.5 (CH-CO), 28.6 (CH₃-CO). Exact Mass (IE): 755.9593 (calcd. 755.9599 for [M]⁺ with (¹⁹³Ir)(⁷⁹Br)); m/z: 658.8 (s, [M-acac]⁺), 579.0 (m, [M-acac-Br]⁺), 499 (w, [M-acac-(Br)₂]⁺). C₂₇H₂₁Br₂IrN₂O₂·H₂O: calcd. C 41.82, H 2.99, N 3.61; found C 42.02, H 2.60, N 3.54.

Synthesis of [Ir(2-(6-p-pyridyl-phenyl)-9,9-dihexylfluorene)₂μ-Cl]₂

Synthetic route is analogous to synthesis of [Ir(2(4-bromophenyl)pyridine)₂μ-Cl]₂. Yield: 66%. ¹H NMR (250 MHz, CDCl₃): δ = 9.45 (d, 3J = 5.4 Hz, 4H; CH-N), 7.95 (d, 3J = 7.2 Hz, 4H; CH-C-C(CH₃)-N), 7.80 (t, 3J = 7.2 Hz, 4H; CH-CH-C-C(CH₃)-N), 7.63 (br, 4H; CH-CH-CH-N), 7.56 (mp, 3J = 6.7 Hz, 4H; CH-C-N), 7.26 (br mp, 20H; ArH), 7.12 (s, 4H; FIH), 7.10 (mp, 4H, FIH), 6.89 (t, 3J = 5.4 Hz, 4H; CH-CH-N), 6.35 (s, 4H; CH-ClIr), 1.7-2.0 (br mp, 16H; CH₂-C), 0.9-1.2 (br mp, 48H, CH₂), 0.78 (br mp, 24H; CH₃), 0.4-0.6 (br, 16H; CH₂).

Synthesis of N (Ir(2-(6-p-pyridyl-phenyl)-9,9-dihexylfluorene)₂(acac))

Synthetic route is analogous to synthesis of M. Yield: 78%. ¹H NMR (250 MHz, CDCl₃): δ = 8.63 (d, 3J = 5.7 Hz, 2H; CH-N), 7.92 (d, 3J = 7.9 Hz, 2H; CH-C-C-N), 7.76 (tr, 3J = 7.2 Hz and 4J = 1.4 Hz, 2H; ArH), 7.64 (d, 3J = 6.0 Hz, 2H; ArH), 7.58 (d, 3J = 5.9 Hz, 2H; ArH), 7.2-7.3 (br mp, 3J = 8.0 Hz, 12H; ArH), 7.1-7.2 (mp, 3J = 7.1 Hz, 4H; ArH), 6.59 (s, 2H, CH-ClIr), 1.8-2.0 (br, 8H; CH₂-C), 0.9-1.2 (br mp, 24H, CH₂), 0.77 (mp, 12H; CH₃), 0.4-0.6 (br, 8H; CH₂). ¹³C{¹H} NMR (500 MHz, CDCl₃): δ = 184.7 (CO), 168.5 (CN), 150.8, 150.6 (C-C-CH₂), 148.4 (CH-N), 144.0 (C-Ir), 141.4, 140.9, 140.4, 139.9 (ArC), 136.7 (CH-CH-CH-N), 126.7, 126.6, 125.8, 124.0, 122.7, 121.4, 121.3, 121.1, 120.0, 119.5, 119.3, 118.4 (CH), 100.4 (CH-CO), 54.9 (C-CH₂), 40.53, 40.48 (C-CH₂), 31.5, 29.8 (CH₂), 28.8 (CH₃-CO), 23.6 (C-CH₂-CH₂), 22.6 (CH₃-CH₂), 14.0, 13.9 (CH₃). Mass (EI) m/z: 1178.3 (w, [M-(C₆H₁₃)]⁺,

1165.2 (s, $[M-(acac)]^+$), 1009 (m, $[M-(C_6H_{13})_3]^+$). $C_{77}H_{87}IrN_2O_2 \cdot H_2O$: calcd. C 72.10, H 6.99, N 2.18; found C 72.40, H 6.87, N 2.27.

Synthesis of O (Ir(2-(6-pyridyl-phenyl)-7-(9,9-dihexylfluorenyl)-9,9-dioctylfluorene)₂(acac)). Synthetic route is analogous to synthesis of N with some minor alterations. The dichlorobridged diiridium complex was prepared in 2-ethoxyethanol and precipitated by adding water. Yield: 59%. 1H NMR (400 MHz, $CDCl_3$): δ = 8.65 (d, 3J = 5.1 Hz, 2H; CH-N), 7.86 (d, 3J = 7.8 Hz, 2H; CH-C-C-N), 7.75-7.58 (mp, 24H; ArH), 7.37-7.16 (mp, 10H; ArH), 6.61 (s, 2H; CH-CH-N), 5.27 (s, 1H; CH-CO), 2.03-1.94 (mp, 16H; C-CH₂), 1.85 (s, 6H, CH₃-CO), 1.32-0.91 (mp, 64H, CH₂), 0.88-0.70 (mp, 40H, CH₂-CH₃). $^{13}C\{^1H\}$ NMR (500 MHz, $CDCl_3$): δ = 184.7 (CO), 168.5 (CN), 151.6, 151.4, 151.00, 150.97 (C-C-CH₂), 148.4 (CH-N), 144.1 (C-Ir), 141.5, 140.8, 140.5, 140.2, 139.6 (ArC), 136.7 (CH-CH-CH-N), 126.9, 126.7, 126.0, 125.9, 124.0, 122.9, 121.5, 121.34, 121.29, 119.8, 119.7, 119.4, 118.4 (CH), 100.8 (CH-CO), 55.1, 55.0 (C-CH₂), 40.6, 40.5, 40.3 (C-CH₂), 31.78, 31.77, 31.4, 30.2, 29.7, 29.3, 29.2 (CH₂), 28.8 (CH₃-CO), 23.7 (C-CH₂-CH₂), 22.58, 22.57, 22.54 (CH₃-CH₂), 14.06, 13.99 (CH₃). Mass (IE) m/z : 2041.4 (s, $[M]^+$), 1942.4 (m, $[M-(acac)]^+$). $C_{135}H_{167}IrN_2O_2$: calcd. C 79.40, H 8.24, N 1.37; found C 79.12, H 8.13, N 1.17.

Synthesis of P (Ir(2-(6-pyridyl-phenyl)-7-(2-(7-[2-(9,9-dihexylfluorenyl)])-9,9-dioctylfluorenyl)-9,9-dioctylfluorene)₂(acac)). Synthetic route is analogous to synthesis of N. Yield: 57%. 1H NMR (400 MHz, $CDCl_3$): δ = 8.65 (d, 3J = 5.4 Hz, 2H; CH-N), 7.94 (d, 3J = 8.3 Hz, 2H; CH-C-C-N), 7.82-7.78 (mp, 8H; ArH), 7.75-7.72 (t, 3J = 8.2 Hz, 4H; ArH), 7.67-7.63 (mp, 22H; ArH), 7.38-7.31 (mp, 8H; ArH), 7.21-7.17 (mp, 4H; ArH), 6.61 (s, 2H; CH-CH-N), 5.28 (s, 1H; CH-CO), 2.11-2.02 (mp, 24H; C-CH₂), 1.85 (s, 6H, CH₃-CO), 1.20-1.07 (mp, 104H, CH₂), 0.82-0.67 (mp, 60H, CH₂-CH₃). $^{13}C\{^1H\}$ NMR (500 MHz, $CDCl_3$): δ = 184.7 (CO), 166.4 (CN), 151.7, 151.6, 151.4, 151.0 (C-C-CH₂), (CH-N not resolved), 143.3 (C-Ir), 140.7, 140.4, 140.3, 139.9 (ArC), (CH-CH-CH-N not resolved), 126.9, 126.7, 126.1, 126.0, 125.9, 122.8, 121.4, 119.9, 119.8, 119.7 (CH), 100.8 (CH-CO), 55.3, 55.12, 55.06 (C-CH₂), 40.5, 40.3 (C-CH₂), 31.7, 31.4, 30.1, 30.0, 29.6, 29.2, 29.1 (CH₂), 28.7 (CH₃-CO), 23.9, 23.7 (C-CH₂-CH₂), 22.53, 22.51 (CH₃-CH₂), 14.0, 13.9 (CH₃). Mass (IE)

m/z: 2819 (m, $[M]^+$), 2719 (w, $[M-(acac)]^+$). $C_{193}H_{247}IrN_2O_2$: calcd. C 82.22, H 8.83, N 0.88; found C 81.93, H 8.77, N 1.10.

Synthesis of ppy-pf8-ppy **Q**: Poly-Suzuki reaction of 9,9-di-n-octylfluorene containing 2-(6-p-pyridyl-phenyl) endgroups on both sides

To a solution of 500 mg (0.78 mmol) of 2,7-di(4,4,5,5,-tetramethyl-1,3,2,-dioxaboronate)-9,9-dioctylfluorene, 405.5 mg (0.740 mmol) of 2,7-dibromo-9,9-dioctylfluorene and 17.1 mg (0.078 mmol) 2-(6-p-pyridyl-phenyl) in 3 ml toluene was added: 4 mg (0.004 mmol) $Pd(PPh_3)_4$ and 2.5 ml of (3.1 mmol) Et_4NOH . The resulting mixture was stirred for 24 h at 100°C. After 3h, 5 ml toluene was added to dissolve the precipitated oligomers or to rise the viscosity of the mixture. After the mixture was cooled down to 20°C the mixture was slowly transferred in a large excess of MeOH, allowing the polymer to precipitate as a white (slightly greenish) fibre-like solid. The solids were collected on a filter, redissolved in 50 ml of toluene and subsequently filtered on a short column of silica. The filtrate was transferred into MeOH resulting in a reprecipitation of the polymer as off-white fibres. The solvents were removed under reduced pressure. 1H NMR (250 MHz, $CDCl_3$): δ = 8.72 (br d, 2H*12%; CH-N), 8.13 (br d, 2H*12%, CH-CH-CH-N), 7.9-7.5 (br, 6H, ArH), 2.3-2.0 (br, 4H, CH_2 -C, 1.3-1.0 (br, 20H, CH_2), 0.9-0.7 (br, 10H, CH_3+CH_2). Mw (NMR): 10.000 (24-mer) GPC: Mp 48586, Mn 18618, Mw 67120, polydispersity 3.6.

Synthesis of ppy-pf8 **E** Suzuki polymerisation of 9,9-di-n-octylfluorene containing 2-(6-p-pyridyl-phenyl) endgroup on one side.

To a solution of 500 mg (0.56 mmol) of 2-(4,4,5,5,-tetramethyl-1,3,2,-dioxaboronate)-7-bromo-9,9-dioctylfluorene and 13.1 mg (0.056 mmol) 2-(6-p-pyridyl-phenyl) in 5 ml toluene was added: 4 mg (0.004 mmol) $Pd(PPh_3)_4$ and 2.5 ml (3.1 mmol) of a Et_4NOH solution in water. The resulting mixture was stirred for 24 h at 110°C. After the mixture was cooled down to 20°C the mixture was slowly transferred in a large excess of MeOH, allowing the polymer to precipitate as a yellow-white solid. The solids were collected on a filter, redissolved in 50 ml of toluene and subsequently filtered on a short column of silica. The filtrate was transferred into MeOH resulting in a reprecipitation of the polymer as off-white fibres. The solvents were removed under reduced pressure. 1H NMR (250 MHz, $CDCl_3$): δ = 8.75 (br d, 2H*12%; CH-

N), 8.13 (br d, 2H*12%, CH-CH-CH-N), 7.9-7.5 (br, 6H, ArH), 2.3-2.0 (br, 4H, CH₂-C, 1.3-1.0 (br, 20H, CH₂), 0.9-0.7 (br, 10H, CH₃+CH₂). Mw (NMR): 5000 (12-mer).

Synthesis of **R**₁; [Ir(acac)(ppy-pf8-ppy)₂Ir(acac)]_n via the iridium complexation of **Q**. A mixture of 200 mg **Q** (containing 0.159 mmol 2-(6-p-pyridyl-phenyl) and 12.5 mg IrCl₃ (containing 0.0353 mmol of iridium) was transferred in a solvent mixture of 5 ml of toluene and 3 ml of 2-ethoxyethanol. The resulting clear brownish mixture was heated to 100°C for 24 h and an orange gel was obtained which further precipitated in a large excess of MeOH. The product was collected on a glass fritted filter, washed with H₂O, MeOH and acetone, and finally the solvents were removed under reduced pressure. Subsequently, a suspension of 161 mg of this product, with 20 mg (0.2 mmol) acetylacetone and 20 mg (0.19 mmol) of Na₂CO₃ in 5 mL of 2-ethoxyethanol and 10 mL of toluene was heated at 90°C for 24 h. After the reaction mixture was cooled down to room temperature the polymer was precipitated in a large excess of MeOH, collected on a filter and washed with H₂O, MeOH and acetone. The polymer was redissolved in 30 mL toluene upon heating at 90°C for 4 h. The resulting mixture was first filtered over glasswool and then over a short silica column after which the polymer was reprecipitated using MeOH. The resulting polymer shows remarkable low solubility in most common solvents; it was found soluble in toluene, dichloromethane and benzene only after lengthy solication. Yield: 161 mg. ¹H NMR (400 MHz, CDCl₃): δ = 7.9-7.5 (br, 6H, ArH), 2.3-2.0 (br, 4H, CH₂-C), 1.3-1.0 (br, 20H, CH₂), 0.9-0.7 (br, 10H, CH₃+CH₂) (signals of iridium complex not resolved). Microanalysis: C 84.58, H 10.18, N 0.42, Ir (AES) 0.65.

Synthesis of **R**₂; [Ir(acac)(ppy-pf8-ppy)₂Ir(acac)]_n via a poly-Suzuki reaction of 2,7-di(4,4,5,5,-tetramethyl-1,3,2,-dioxaboronate)-9,9-di-n-octylfluorene, 2,7-dibromo-9,9-dioctylfluorene and N (Ir(2(4-bromophenyl)pyridine)₂acac)

A solution of 500 mg (0.779 mmol) 2,7-di(4,4,5,5,-tetramethyl-1,3,2,-dioxaboronate)-9,9-dioctylfluorene, 405.5 mg (0.740 mmol) of 2,7-dibromo-9,9-dioctylfluorene, 29.5 mg (0.0389 mmol) of **M** (Ir(2(4-bromophenyl)pyridine)₂acac) and 4 mg (0.004 mmol) of (Pd(PPh₃)₄) in 10 mL of toluene was heated to 90°C and 2.5 mL (3.1 mmol) of Et₄NOH solution in water was added. The resulting two-phase mixture was stirred for 18 h at 100°C and formed a brown-orange gel. The gel was precipitated in MeOH and

washed with MeOH. The polymer was redissolved in 30 mL of dichloromethane and the resulting mixture was filtered using glass wool and subsequently filtered over a short silica column. The resulting polymer is soluble in toluene and dichloromethane after lengthy sonication, and gives high viscosity. ^1H NMR (400 MHz, C_6D_6): δ = 8.2-7.6 (br, 6H, ArH), 2.6-1.9 (br, 4H, $\text{CH}_2\text{-C}$), 1.6-0.7 (br, 30H, $\text{CH}_2\text{+CH}_3$) (spectrum is very broad, signals of iridium complex not resolved). GPC: Mp 50643, Mn 28146, Mw 62638, polydispersity 2.2. Microanalysis: C 84.28, H 9.71, N 0.37, Ir (AES) 2.56.

Synthesis of S_1 ; $[\text{Ir}(\text{ppy-pf8})_2(\text{acac})]$ via iridium complexation of **E**

A mixture of 150 mg **E** (containing 0.0367 mmol 2-(6-pyridyl-phenyl)) and 5.2 mg IrCl_3 (containing 0.0147 mmol of iridium) was transferred in a solvent mixture of 10 ml of toluene and 10 ml of 2-ethoxyethanol. The resulting clear yellow-orange mixture was heated to 110°C for 24 h after which the product was precipitated in a large excess of MeOH. The product was collected on a glass fritted filter, washed with MeOH and hexane, and finally the solvents were removed under reduced pressure. Subsequently, a clear solution the product and 20 mg (0.2 mmol) acetylacetone and 20 mg (0.19 mmol) of Na_2CO_3 in 8 mL of 2-ethoxyethanol and 2 mL of toluene was heated at 110°C for 2 h. Upon cooling to 0°C a yellow-orange precipitate was formed that was collected on a glass fritted filter, washed with H_2O and MeOH and the solvents were removed under reduced pressure. The product was filtered over a short silica column using toluene as the solvent. Yield: 100 mg. ^1H NMR (250 MHz, CDCl_3): δ = 8.65 (br, 1H*1%; CH-N), 7.9-7.5 (br, 6H, ArH), 6.62 (br, 1H*1%, CH-CHN), 5.30 (br, 0.5H*1%; CH-CO), 2.3-2.0 (br, 4H, $\text{CH}_2\text{-C}$), 1.82 (br, 3H*1%, $\text{CH}_3\text{-CO}$), 1.3-1.0 (br, 20H, CH_2), 0.9-0.7 (br, 10H, $\text{CH}_3\text{+CH}_2$). Microanalysis: C 83.09, H 9.61, N 0.39.

Synthesis of S_2 : $[\text{Ir}(\text{ppy-pf8})_2(\text{acac})]$ via a poly-Suzuki reaction using **C** 2-(4,4,5,5-tetramethyl-1,3,2-dioxaborolane)-7-bromo-9,9-dioctylfluorene and **M** ($\text{Ir}(\text{2(4-bromophenyl)pyridine})_2\text{acac}$)

A solution of 415 mg (0.697 mmol) of **C**, 26.4 mg (0.0349 mmol) of **N**, 2 mg (0.008 mmol) of $\text{Pd}(\text{acetate})$ and 4 mg (0.016 mmol) of PCy_3 was heated to 90°C and 2.5 mL (3.1 mmol) of a Et_4NOH solution in water was added. The resulting mixture was

stirred at 110°C for 20 h, resulting in a slightly viscous clear red solution. The organic layer was dropped out in a large excess of MeOH, which resulted in the precipitation of an orange polymer. This was washed with water, MeOH and acetone and filtered over a short silica column using toluene as the eluent. The resulting solution was concentrated to 2 ml and 100 mg of Na₂CO₃, 1 mL of acacH and 8 mL of 2-ethoxyethanol were added. The reaction mixture was stirred for 2 h at 110°C and subsequently cooled down to room temperature. 2 mL of MeOH was added on which the product precipitated as an orange solid. The product was isolated on a glass fritted filter, washed with H₂O and MeOH and the solvents were removed under reduced pressure. Yield: 150 mg (55%). δ = 8.7 (d, J = 4.6 Hz, 1H*10%; CH-N), 7.9-7.5 (br, 6H; ArH), 7.4-7.1 (br, ArH), 6.6 (s, 1H*10%; CH-CH-N), 5.3 (s, 0.5H*10%; CH-CO), 2.3-2.0 (br, 4H, CH₂-C), 1.9 (3H*10%; CH₃-CO), 1.3-1.0 (br, 20H, CH₂), 0.9-0.7 (br, 10H, CH₃+CH₂). Microanalysis: C 82.78, H 9.44, N 0.25.

S₃ was synthesised analogues to S₂ but using a solution of 415 mg (0.697 mmol) of C and 13.2 mg (0.0349 mmol) of N. Yield: 230 mg (85%). δ = 8.7 (d, J = 5.0 Hz, 1H*3%; CH-N), 7.9-7.5 (br, 6H; ArH), 7.4-7.1 (br, ArH), 6.6 (s, 1H*3%; CH-CH-N), 5.3 (s, 0.5H*3%; CH-CO), 2.3-2.0 (br, 4H, CH₂-C), 1.9 (3H*3%; CH₃-CO), 1.3-1.0 (br, 20H, CH₂), 0.9-0.7 (br, 10H, CH₃+CH₂). Microanalysis: C 87.10, H 10.02, N 0.20.

LED fabrication and measurements

The electroluminescent devices have been fabricated as described below. An indium-tin oxide (ITO) patterned and cleaned glass substrate was obtained from Cambridge Display Technology Limited. A hole-injection layer, Poly(3,4-ethylenedioxythiophene):poly(styrene sulfonic acid) (PEDOT:PSS, obtained from Baytron as an aqueous solution and used without further pretreatment), was spin-coated (2000 rpm, 30 seconds) onto the ITO. The PEDOT:PSS layer was dried on a hotplate (130 °C) for 5 minutes. Next, a hole-transport layer of polyvinylcarbazole (PVK) was spin-coated, from a 3 mg/mL solution in dichloromethane, on top of the PEDOT:PSS layer. Subsequently, the light-emitting polymer was spin-coated (from a 10-15 mg/mL solution in toluene) on top of this layer. Then, the device was transferred to a high-

vacuum (1×10^{-5} mbar) deposit unit and set with a shadow mask. A cathode metal, calcium, was deposited on the organic layers through the shadow mask, followed by deposition of aluminium. The size of each pixel is $4 \text{ mm} \times 3.2 \text{ mm}$. The device was finally encapsulated using Araldite epoxy resin and covered with a glass slide. Typical characteristics of the device layers include: ITO (190 nm, sheet resistance $< 10 \Omega/\square$) / PEDOT:PSS (50 nm) / PVK (45 nm) / light-emitting polymer (70 nm) / Ca (40 nm) / Al (200 nm). The devices were examined by applying a direct current bias (0 to 25 V) to the electrodes. The electroluminescent characteristics were measured with a Hewlett-Packard E3631A DC Power Supply, Keithley 2000 Digital Multimeter, Topcon BM-9M Luminance Meter, and Aminco-Bowman Series 2 Fluorescence Spectrometer.

Results and Discussion

Synthesis of iridium(ppy)-fluorenyl hybrid systems

The 4-position of 2-phenylpyridine (ppy) was functionalised with a series of 9,9-dialkylfluorene segments including one, two and three subsequent fluorenyl substituents (Scheme 1).

Scheme 1: Synthetic pathway to 9,9-dialkylfluorene substituted 2-phenylpyridine ligands

(2(4-bromophenyl)pyridine) (A) was synthesised via a coupling of pyridine to parabromoaniline using the Gomberg-Hey reaction.⁵ Monofluorenyl functionalised ppy was obtained by a Suzuki coupling of A with 2-(4,4,5,5 tetramethyl-1,3,2-dioxaborolane)-9,9-dihexylfluorene to obtain (2-(6-p-pyridyl-phenyl)-9,9-dihexylfluorene) (D). The key building block of our stepwise synthesis; 2-(4,4,5,5 tetramethyl-1,3,2-dioxaborolane)-7-trimethylsilyl-9,9-dioctylfluorene (B) was synthesised in two steps from 2,7-dibromo-9,9-dioctylfluorene via a monolithiation using nBuLi, quenched with TMSCl followed by a second lithiation using tBuLi, that was further reacted with 2-isopropoxy-4,4,5,5 tetramethyl-1,3,2-dioxaborolane to obtain B. 4-Difluorenyl functionalised ppy was obtained in two steps from the Suzuki

product of **A** and **B** via the deprotection of the TMS group using ICl (which yields 2-(6-p-pyridyl-phenyl)-7-tetramethyl-1,3,2-dioxaborolane)-9,9-dihexylfluorene to obtain 2-(6-p-pyridyl-phenyl)-7-(9,9-dihexylfluorenyl)-9,9-dioctylfluorene (**H**).⁶ 4-Trifluorenyl functionalised ppy was obtained from the Suzuki product of **G** and 2-(4,4,5,5 tetramethyl-1,3,2-dioxaborolane)-9,9-dihexylfluorene via the same deprotection/Suzuki reaction sequence to obtain 2-(6-p-pyridyl-phenyl)-7-(2-(7-[2-(9,9-dihexylfluorenyl)]))-9,9-dioctylfluorenyl)-9,9-dioctylfluorene (**K**).

Scheme 2: Synthesis of 4-fluorenyl functionalised ppy ligands

Oligofluorene-bis-ppy was synthesised using a $\text{Pd}(\text{PPh}_3)_4$ catalysed poly-Suzuki reaction using **A**, 2,7-di(4,4,5,5-tetramethyl-1,3,2-dioxaboronate)-9,9-dioctylfluorene and 2,7-dibromo-9,9-dioctylfluorene as the building blocks to yield ppy-pf8-ppy (**Q**) (Scheme 2).⁷ Oligofluorene-mono-ppy was obtained from the same coupling reaction using **A** and 2-(4,4,5,5 tetramethyl-1,3,2-dioxaborolane)-7-bromo-9,9-dioctylfluorene (**C**) as the building blocks to obtain ppy-pf8 (**E**) (Scheme 2).

Bis-cyclometalated iridium complexes of the type $\text{Ir}(\text{ppy-X})_2(\text{acac})$ were synthesised according to a two-step synthetic route that was recently reported by Lamansky et al.⁸ In the first step, $\text{IrCl}_3 \cdot n\text{H}_2\text{O}$ is reacted with an excess of the ppy containing ligand to yield the chloro-bridged dimer $[\text{Ir}(\text{ppy-X})_2(-\mu\text{-Cl})]_2$ (Scheme 3)

Scheme 3: Synthesis pathway to bis-CN-cyclometalated iridium complexes

The chloro-bridges iridium dimers are subsequently converted to phosphorescent monomeric acetylacetonate complexes. $(\text{Ir}(2(4\text{-bromophenyl})\text{pyridine})_2\text{acac})$ (**M**), $(\text{Ir}(2-(6\text{-p-pyridyl-phenyl})-9,9\text{-dihexylfluorene})_2(\text{acac}))$ (**N**), $(\text{Ir}(2-(6\text{-p-pyridyl-phenyl})-7-(9,9\text{-dihexylfluorenyl})-9,9\text{-dioctylfluorene})_2(\text{acac}))$ (**O**) and $\text{Ir}(2-(6\text{-p-pyridyl-phenyl})-7-(2-(7-[2-(9,9\text{-dihexylfluorenyl}]))) - 9,9\text{-dioctylfluorenyl})-9,9\text{-dioctylfluorene})_2(\text{acac}))$ (**P**) were all synthesised accordingly (Scheme 3).

The synthesis of iridium(ppy)-oligofluorenyl systems was attempted via two approaches (Scheme 4).

Scheme 4: Synthetic pathways towards Ir(ppy)-oligofluorenyl systems

Analogous to the model compounds, both **Q** and **E** were exposed to the two-step synthetic pathway via the chloro-bridged iridium dimer towards the acetylacetonate containing monomer. This seemed troublesome using **Q**, since a full conversion to the chloro-bridged iridium dimer was frustrated by gelation of the reaction mixture. AES analyses on the targeted product; $[\text{Ir}(\text{acac})(\text{ppy-pf8-ppy})_2]_n$ (**R**₁) revealed that the iridium content was only 0.65% (objected: 4%). Moreover, the resulting polymer did only dissolve after extensive sonication. Upon complexation of **E** the gel-formation was completely avoided due to the prevention of network formation of the dichloro-bridged intermediate. Hence, the product of this two-step reaction sequence; $[\text{Ir}(\text{ppy-pf8})_2(\text{acac})]$ (**S**₁) is readily soluble in dichloromethane and toluene. From NMR it was concluded that 1% $\text{Ir}(\text{ppy})_2(\text{acac})$ segments are present (while 4% was objected).

Alternatively, complex **M** was applied in a poly-Suzuki reaction using 2,7-di(4,4,5,5,-tetramethyl-1,3,2,-dioxaboronate)-9,9-dioctylfluorene and 2,7-dibromo-9,9-dioctylfluorene as the building blocks to yield $[\text{Ir}(\text{acac})(\text{ppy-pf8-ppy})_2]_n$ (**R**₂). Also via this pathway the synthesis of **R** gave rise to gel formation during the reaction and caused solubility problems in the manufacturing of it. This method did however give rise to an increased inbuild of $\text{Ir}(\text{ppy})(\text{acac})$ segments compared to approach **R**₁; in this case the iridium contents was 2.6%. The most successful iridium(ppy)-oligofluorenyl system was obtained via a poly-Suzuki reaction using **C** and **M**. The resulting product $[\text{Ir}(\text{ppy-pf8})_2(\text{acac})]$ (**S**₂) was highly soluble in dichloromethane and toluene and contained 5% of the bis-cyclometalated iridium complex. A subsequent reaction with acacH was needed to restore the acac-anion back on the complex.

Photophysical properties of $\text{Ir}(\text{ppy-X})_2(\text{acac})$ complexes

Absorption and photoluminescence (PL) spectra were recorded of the series of cyclometalated iridium complexes to investigate the influence of the fluorenyl-substitution on the 4-position of the ppy (Figure 1).

Figure 1: Absorption and PL-spectra of $\text{Ir}(\text{ppy-X})_2(\text{acac})$ complexes

The parent compound **L** shows an *absorption* band and 260 nm while it emits at 507 nm. When a monofluorenyl-segment is placed at the 4 position of the ppy (complex **N**), the absorption maximum was observed at 338 nm with a second, minor maximum at 260 nm. Clearly, complex **N** absorbs both from the π system as well as from the d-orbitals of the iridium complex. Further extension of the system of the ligands with subsequent fluorenyl units (via complexes **O** and **P**) resulted in a red shift of the absorption maximum from the π system of the ligand accompanied by a relative decrease of the absorption intensity originating from the transition metal. Complex **P** shows a maximum absorption at 376 nm which is only 8 nm lower than where polyfluorene absorbs. For this complex, the absorption from the transition metal is only observed as a minor absorption peak at 260 nm. In comparison; when the 4-position of the ppy was substituted by a bromine, it had no impact on the absorption spectrum at all.

Interestingly, compared to the unsubstituted complex, the *PL-spectrum* of complex **N** (one fluorenyl) is red shifted with respect to **L** by 30 nm, to 552 nm. From this it can be concluded that the triplet wavefunction in this system is not confined to the Ir(ppy) unit but extended over the complete system. A second fluorenyl-segment (complex **O**) gave rise to a further red shift of 9 nm (to 561 nm) and a third (**P**) provided a PL-maximum of 566 nm.

In general it appears that the influence of the repeat units of the fluorenyl-chain on the wavefunction is gradually decreasing when positioned further away from the phosphor, but that at least the first three fluorenyl units influence the triplet wavefunction (in our case by causing a total red shift of 44 nm).

EL-characteristics were investigated by manufacturing OLED's (organic light emitting diode) from these model compounds. The OLED device consists of a basic three layered structure sandwiched between indium tin oxide and a calcium electrode; ITO/PEDOT-PSS/PVK/Emitting layer/Ca/Al. The use of PEDOT-PSS is commonly known as a hole-injection layer, the polyvinylcarbazole layer (PVK) is added as a hole transport layer and was found to generally smoothen the device characteristics. The series of **L**, **N**, **O** and **P** were spincoated in this structure from a toluene solution. The EL-spectra of these devices roughly show the same trend as the PL-spectra. In this series, the emission maximum shifts red from 529 nm for **L**, via 561 nm (**N**), to

567 nm for **O** and **P**. Apart from a little red-shift all of the EL-spectra of these compounds closely resemble their PL-spectra. Not a trace of fluorenyl originating singlet emission from these systems was observed.

Figure 2: EL spectra of Ir(ppy-X)₂(acac) (inset shows the device structure)

Table 1: EL-characteristics of Ir(ppy-X)₂(acac)

System	Lum. max (cd/m ²)	turn on voltage (V)	CIE X	CIE Y	Efficien cy (cd/A)	ELQY (%)	Colour
L	60	4.0	0.37	0.59	$2.5 \cdot 10^{-3}$	$7.4 \cdot 10^{-4}$	yellowish h- green
N	43	5.0	0.48	0.51	$9.9 \cdot 10^{-3}$	$3.3 \cdot 10^{-3}$	greenish yellow
O	400	5.5	0.48	0.51	$6.9 \cdot 10^{-2}$	$2.2 \cdot 10^{-2}$	yellow
P	1400	6.0	0.46	0.53	$7.3 \cdot 10^{-2}$	$7.3 \cdot 10^{-2}$	greenish yellow

The device properties improve substantially on extending the ligand system with fluorenyl segments (Table 1 and Figure 3). The maximum brightness increased from 60 (for **L**), via 43 for **N**, 400 for **O** to 1400 for **P**, while the quantum yield improved two orders of magnitude from $7.4 \cdot 10^{-4}$ to $7.3 \cdot 10^{-2}$.⁹ The turn on voltage went up slightly upon increasing the percentage of fluorene.

From the experiments described above it was concluded that the covalent bonding of fluorenyl segments directly to the system of the ligand system of this type of phosphorescent complexes is beneficial for its ability to harvest triplets. This phenomenon is not explained solely by the improved film forming capabilities of the higher fluorenyl substituted phosphors. We suggest that upon increasing the system of the phosphor, more uniform films are obtained with a more homogeneous distribution of the iridium(ppy) species, avoiding clustering which can give rise to

triplet annihilation. The size of the system of the phosphors might also be of importance in the efficiency of capturing charges.

Figure 3: I-L-V curves of Ir(ppy-X)₂(acac)

The results prompted us to investigate this type of phosphors in a polyfluorene matrix. PL measurements of blends of complex L and N in a 9,9-dioctyl-polyfluorene (PF8) matrix revealed that singlet emission from the polyfluorene was significantly quenched by both phosphors, but that the triplet emission (at room temperature under aerobic conditions) from both systems was very low (Figure 4). Exciton *Föster* transport from the S₁ of the polymer to the T₁ seems to be efficient, but a different mechanism (triplet annihilation, or *Dexter* transfer of triplets from the phosphor back to the polymer) seems responsible for a major quenching of excitons on the phosphor.

Figure 4: PL spectra of blends of complex N in PF8. Inset; triplet signal of blends of 5.0 mol % of complex L and N in PF8

The EL spectrum of L in PF8 showed a marked resemblance with that of undoped PF8, but the emission maximum was shifted from 516 to 533 (note that the emission maximum of 516 nm of PF8 is caused by excimer formation due to aggregation effects) (Figure 5). It seems that this device emits both from the singlet excited state of the polymer (a maximum at 414 nm) as well as from the triplet excitons on the transition metal complex (533 nm). The blend of complex N in PF8 showed nearly only triplet emission. The emission maximum at 559 nm is clearly originating from the phosphor (552 nm was observed for the neat complex; Figure 2) whereas the singlet peak at 443 nm is very small. We owe this difference between complex L and

Figure 5: EL spectra of Ir(ppy-X)₂(acac) in polyfluorene

N to the difference in T_1 -level. From the emission spectra it was estimated that the T_1 -level of L is about 2.4 eV, and that of complex N is 2.2 eV. On comparing this with the T_1 level of PF8, which is recently calculated to be about 2.30 eV¹⁰ it can be predicted that in the case of L part of the triplets, captured on the phosphor, will hop to the T_1 -level (*Dexter* transfer) of the polymer and decay non-radiatively. Hence triplet emission is frustrated by this process. In the case of complex N such a process is less likely since the estimated T_1 -level of the polymer is higher than that of the phosphor.

Table 2: EL-data on blends of Ir(ppy-X)₂(acac) in polyfluorene

System	Lum. max (cd/m ²)	turn on voltage (V)	CIE X	CIE Y	Efficien cy (cd/A)	ELQY (%)	Colour
L in PF8	225	5.5	0.35	0.48	0.16	7.6*10 ⁻²	yellowish- green
N in PF8	190	7.5	0.44	0.52	0.15	5.6*10 ⁻²	greenish yellow

The enhanced triplet to singlet ratio of complex N in PF8, compared to the blend of L, doesn't give rise to a significant change in device efficiency (5.6*10⁻² vs 7.6*10⁻², Table 2).

Polyfluorenyl functionalised phosphors

The concept of fluorenyl substituted phosphors was taken one step further by implimentating the bis-cyclometalated iridium phosphors in the main chain of polyfluorene. The PL-spectra of these hybrid systems (**R**₁, **R**₂, **S**₁, and **S**₂) are depicted in Figure 6.

Figure 6: PL-spectra of Ir(ppy-X)₂(acac) incorporated polyfluorenes

All systems show singlet emission (with maxima at 422 and 450 nm) originating from the polyfluorene part of the molecule. In **R**₁ and **R**₂ no triplet emission was observed.

This, together with the NMR-spectra, strongly suggests that in these systems the $\text{Ir(ppy-X)}_2(\text{acac})$ phosphor could not be adequately implemented in the polymer chain due to gelation problems during the synthesis. In this respect systems S_1 and S_2 were more promising since in these materials, only one phosphor can be built in every molecule (Scheme 4) leading to better defined and applicable (soluble) systems. Both S_1 and S_2 showed triplet PL-emission at 596 nm.

The EL-spectra of the novel hybrid system, depicted in Figure 7, again show the marked differences between the R and S products.

Figure 7: EL-spectra of polyfluorene incorporated iridium-complexes

The spectra of R_1 and R_2 are very broad and show a lot of singlet and singlet-excimer emission, caused by aggregation (the low solubility of these compounds gave rise to increased aggregation, compared to normal polyfluorene). R_1 doesn't show any triplet emission at all, while a shoulder at ~ 604 nm could indicate some triplet emission from the device fabricated from system R_2 . The device, containing system S_1 was found to emit singlets and triplets in the ratio 1:3 and S_2 yielded a singlet to triplet ratio of 1:23. This resulted in pink emission (according to the CIE-coordinates $X = 0.42$ and $Y = 0.34$) for S_1 systems whitish orange emission for S_2 ($X = 0.47$ and $Y = 0.43$) (Table 3).

Figure 8: EL-spectra of S_2 at increasing driving voltage, going from 8.5V (bottom) up in steps of 0.5V to 12.5V (top).

The singlet to triplet ratio was found to increase upon increasing the driving voltage (Figure 8). This is explained by an increase in triplet-triplet annihilation at higher driving voltages. The observed increase in excimer formation at increased driving voltages is a wellknown phenomenon in polymer LED's.

Table 3: EL-data on polyfluorenyl-iridium(ppy) complexes

System	Lum. max	turn on voltage	CIE X	CIE Y	Efficiency	ELQY (%)	Colour
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	(cd/m ²)	(V)	(cd/A)				
R ₁	10	8.5	0.29	0.45	7.0*10 ⁻²	2.6*10 ⁻²	yellowish h-green
R ₂	60	8.5	0.33	0.44	0.13	7.1*10 ⁻²	yellowish h- green
S ₁	42	9.5	0.42	0.34	7.6*10 ⁻²	3.6*10 ⁻²	pink
S ₂	163	7.0	0.47	0.43	0.19	7.9*10 ⁻²	whitish- orange
S ₃	304	6.0	0.42	0.40	0.18	7.9*10 ⁻²	whitish- orange

The EL-device characteristics of the hybrid systems are summarized in Table 3. As observed from the EL-spectra the R samples give yellowish-green emission, originating from singlet- and singlet-excimer emission at low brightness (10 cd/m² for R₁ and 60 cd/m² for R₂) at a quantum yield of 2.6*10⁻² and 7.1*10⁻² respectively. The better defined S samples emit mainly from the triplet state and a little from the singlet excited state. S₁ gives pink emission (CIE-X 0.42, CIE-Y 0.34) with a maximum brightness of 42 cd/m² with a quantum yield of 3.6*10⁻². S₂ and S₃ both yield whitish orange emission at a quantum yield of 7.9*10⁻². The relative longer polyfluorene substituents of S₃ (30 fluorenyl units versus 10 for S₂) gives rise to an increase of the maximum brightness from 163 to 304 cd/m².

Conclusion

A series of polyfluorenyl-phosphor composite materials has been studied that give bright phosphorescence in solution-processed devices. The addition of fluorene substituents to the ligands of the small molecule Ir(ppy)₂(acac) yielded materials that are readily spin-coated from solution of blends in fluorene and, more importantly, also as neat materials. The fluorenyl substituents were found to give rise to a major improvement in EL-luminance brightness and in quantum efficiency. A beneficial effect of the substituents was observed in the improved complementation of the T₁-

energy level of the phosphor to that of polyfluorene in blends. OLED's prepared from polyfluorenyl-phosphor composites were reported to yield bright triplet emitting devices.

The novel *phosphors* are hybrid compounds of a bis cyclometalated iridium-2-phenylpyridinato complex (Ir(ppy)_2) triplet emitter, and polyfluorene, an efficient singlet emitting polymer. The series consists of $\text{Ir(ppy)}_2(\text{acac})$ complexes having fluorenyl substituents on the 4-position of the ppy ligand. In the series, from non substituted complex $\text{Ir(ppy)}_2(\text{acac})$ (L) via monofluorenyl substituted complex [$\text{Ir(2-(6-p-pyridyl-phenyl)-9,9-dihexylfluorene)}_2(\text{acac})$] (N) and the difluorenyl [$\text{Ir(2-(6-p-pyridyl-phenyl)-7-(9,9-dihexylfluorenyl)-9,9-dioctylfluorene)}_2(\text{acac})$] (O) to the trifluorenyl [$\text{Ir(2-(6-p-pyridyl-phenyl)-7-(2-(7-[2-(9,9-dihexylfluorenyl)])))-9,9-dioctylfluorenyl)-9,9-dioctylfluorene)}_2(\text{acac})$] (P) a major red-shift was observed in the PL-spectra of these complexes from 522 nm, via 552 nm and 561 to 566 nm. The polyfluorenyl substituted complex [$\text{Ir(ppy-pf8)}_2(\text{acac})$] (S_2) also showed PL triplet emission at 566 nm, along with minor singlet emission at 445 nm. The red-shift of complex N with respect to L of 30 nm was found sufficient to complement the T_1 energy level of the phosphor to that of polyfluorene; the EL spectrum of a blend of N in this polymer shows mainly triplet emission at 559 nm while the blend, containing complex L (λ_{max} at 535 nm), still shows substantial singlet emission at 441 nm. The stepwise elongation of the ppy ligand with fluorenyl segments resulted in a major increase in EL-brightness (from 60 for L to 1400 cd/m^2 for P) and in EL-quantum efficiency (from 7.4×10^{-4} to 7.3×10^{-2}). Yellowish orange EL emission was obtained from S_2 (estimated fluorenyl length; 10 units) with a triplet to singlet ratio of 23:1 at a driving voltage of 7V, with a quantum efficiency of 7.9×10^{-2} and a maximum brightness of 163 cd/m^2 . Upon increasing the length of the polyfluorene substituent to 30 (S_3) yielded whitish orange emission having a triplet to singlet ratio of 7:1 (at 6V), with a quantum efficiency of 7.9×10^{-2} and a maximum brightness of 304 cd/m^2 .

CLAIMS:

1. A material capable of luminescence comprising:
a polymer or oligomer; and
an organometallic group
characterised in that the organometallic group is covalently bound to the polymer or oligomer and the nature, location and/or proportion of the polymer or oligomer and of the organometallic group in the material are selected so that the luminescence predominantly is phosphorescence.
2. A material according to claim 1 wherein the luminescence is electroluminescence.
3. A material according to claim 1 or claim 2 wherein the organometallic is conjugatively bound to the polymer or oligomer.
4. A material according to any one of the preceding claims wherein the polymer or oligomer is semiconducting.
5. A material according to claim 4 wherein the polymer or oligomer is capable predominately of fluorescence in the absence of the organometallic.
6. A material according to claim 5, wherein the polymer or oligomer comprises an aryl or heteroaryl repeat unit.
7. A material according to claim 6, wherein the aryl or heteroaryl repeat unit comprises a group selected from the group consisting of 2,7-linked 9,9 disubstituted fluorene, , a p-linked dialkyl phenylene, a p-linked disubstituted

phenylene, a phenylene vinylene, a 2,5-linked benzothiadiazole, a 2,5-linked substituted benzothiadiazole, a 2,5-linked disubstituted benzothiadiazole, a 2,5-linked substituted or unsubstituted thiophene or a triarylamine.

8. A material according to any one of the preceding claims, wherein the organometallic contains a transition metal.

9. A material according to claim 8, wherein the organometallic contains a precious metal.

10. A material according to any one of the preceding claims wherein material comprises organometallic in an amount in the range from 1 to 10 % by weight.

11. A material according to any one of the preceding claims wherein the organometallic is pendent from the backbone of the polymer or oligomer.

12. A material according to any one of claims 1 to 10 wherein the organometallic forms a part of the backbone of the polymer or oligomer.

13. A material according to claim 12, wherein the organometallic is located at the end of the polymer or oligomer backbone.

14. A material according to claim 12 or claim 13 wherein the organometallic contains an aryl or heteroaryl group.

15. A material according to claim 14, wherein the aryl or heteroaryl group comprises a group selected from the group

consisting of 2,7-linked 9,9 disubstituted fluorene, , a p-linked dialkyl phenylene, a p-linked disubstituted phenylene, a phenylene vinylene, a 2,5-linked benzothiadiazole, a 2,5-linked substituted benzothiadiazole, a 2,5-linked disubstituted benzothiadiazole, a 2,5-linked substituted or unsubstituted thiophene or a triarylamine.

16. Use of a material according to any one of the preceding claims as a component in an optical device.

17. Use of a material according to claim 16, wherein the optical device comprises an electroluminescent device.

18. An optical device or a component therefor, which comprises a substrate and a material as defined in any one of claims 1 to 15 supported on the substrate.

19. An optical device or a component therefor according to claim 18, wherein the optical device comprises an electroluminescent device.

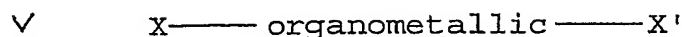
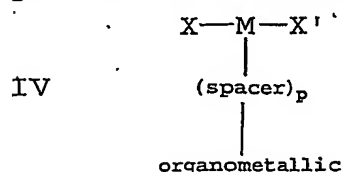
20. An optical device according to claim 19, wherein the electroluminescent device comprises:

- a first charge carrier injecting layer for injecting positive charge carriers;

- a second charge carrier injecting layer for injecting negative charge carriers;

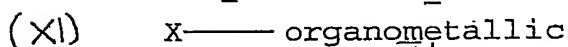
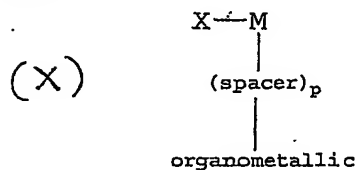
- a light-emissive layer located between the charge carrier injecting layers for generating light and comprising a material as defined in any one of claims 1 to 15.

21. A monomer for use in a polymerisation reaction having a general formula as shown in IV, V or VI below:



where X and X' each is a reactive groups independently selected from the group consisting of a halide group, a boronic acid group, a boronic ester group and a borane group; p o M is a group comprising an aryl or heteroaryl group and L is a ligand capable of forming a complex with a metal when treated with a metal-complexing reagent, preferably the organometallic does not comprise Ru.

22. An end-capping reagent for use in a polymerisation reaction having a general formula as shown in formula X, XI or XII:



where L is a ligand capable of forming a complex with a metal when treated with a metal-complexing reagent; and X is a reactive group selected from the group consisting of a halide

group, a boronic acid group, a boronic ester group and a borane group.

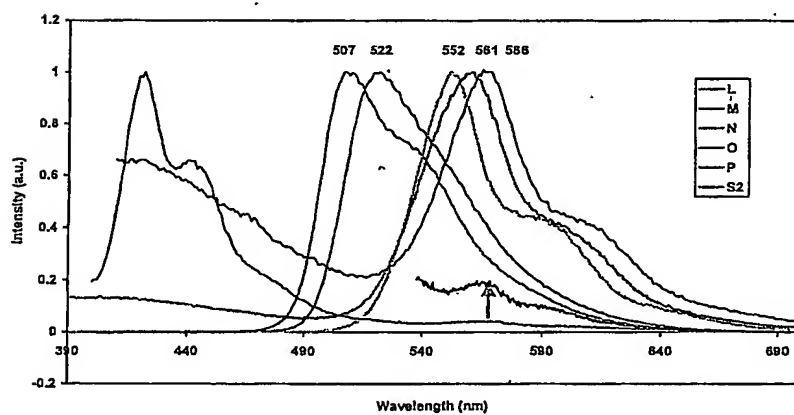
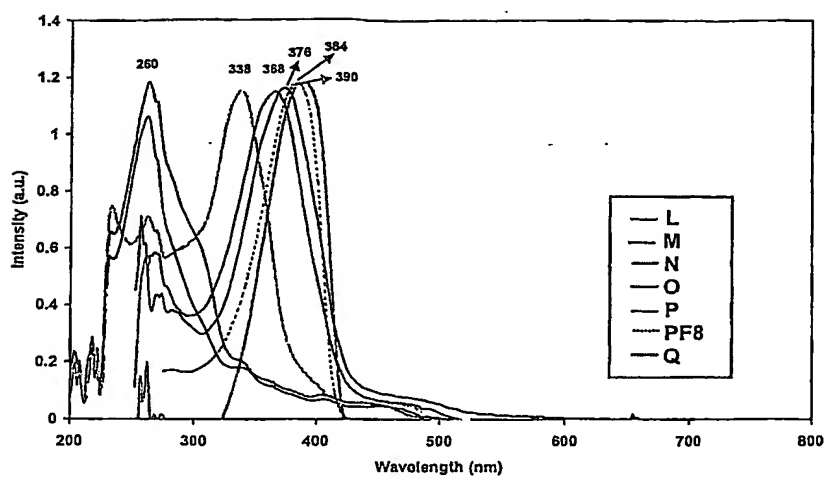
23. A process for preparing a material as defined in claim 13 which comprises:

- (a) reacting monomers to form a polymer or oligomer wherein each monomer has at least two reactive groups selected from the group consisting of a halide group, a boronic acid group, a boronic ester group and a borane group and each monomer comprises an aryl or heteroaryl group; and
- (b) terminating the polymer or oligomer formed in step (a) using an end-capping reagent, said end-capping reagent comprising one reactive group selected from the group consisting of a halide group, a boronic acid group, a boronic ester group and a borane group and either (i) containing an organometallic as defined in claim 1 or claim 9 or (ii) being capable of forming a complex with a metal when treated with a metal-complexing reagent; and
- (c) where the end-capping reagent is as defined in (ii), treating the terminated polymer or oligomer from step (b) with a metal-complexing reagent.

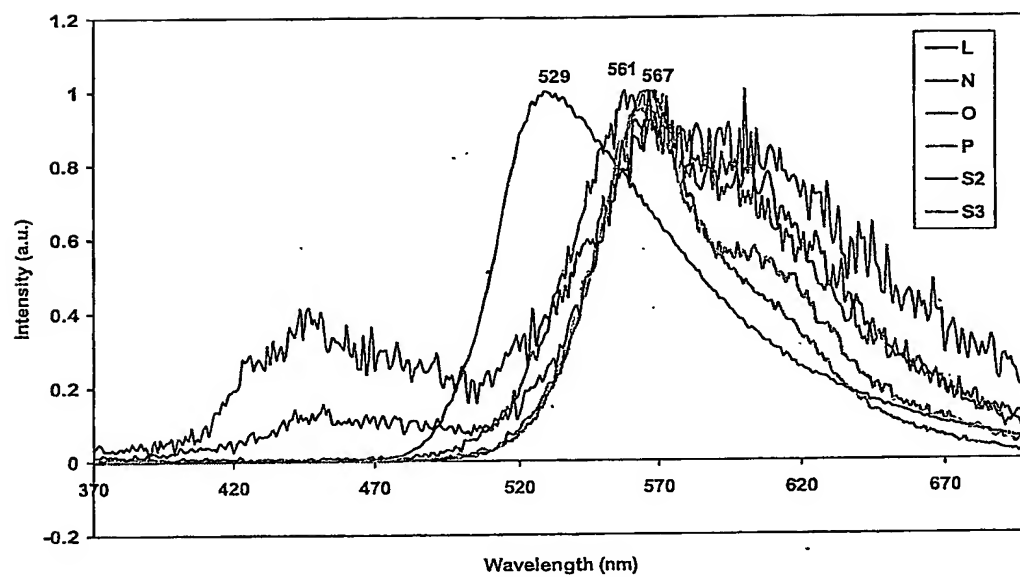
24. A process for preparing a material as defined in any one of claims 1 to 12 which includes reacting at least one first monomer with a plurality of second monomers which are different to the first monomer to form a polymer or oligomer;

wherein each monomer comprises an aryl or heteroaryl group and has at least two reactive groups selected from the group consisting of a halide group, a boronic acid group, a boronic ester group and a borane group; and wherein the first monomer either (i) contains an organometallic or (ii) is capable of forming a complex with a metal when treated with a metal-complexing reagent; and

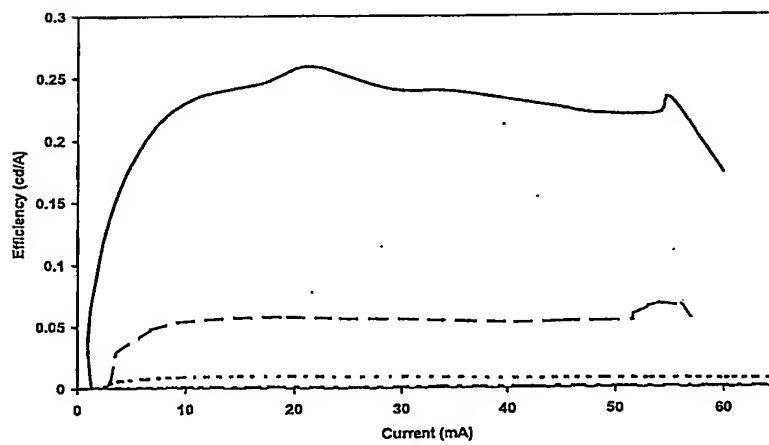
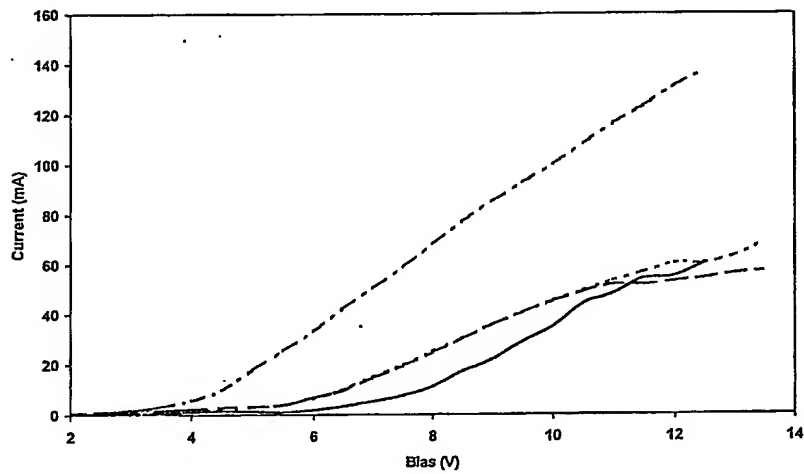
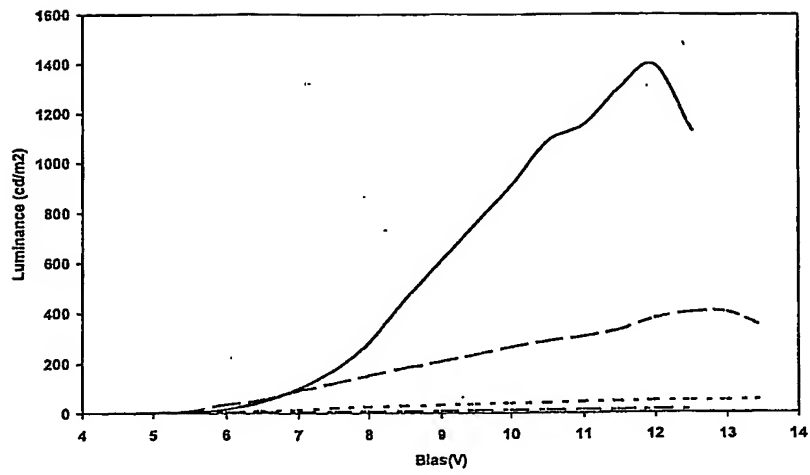
(c) where the first monomer is as defined in (ii), treating the polymer or oligomer from step (b) with a metal-complexing reagent.



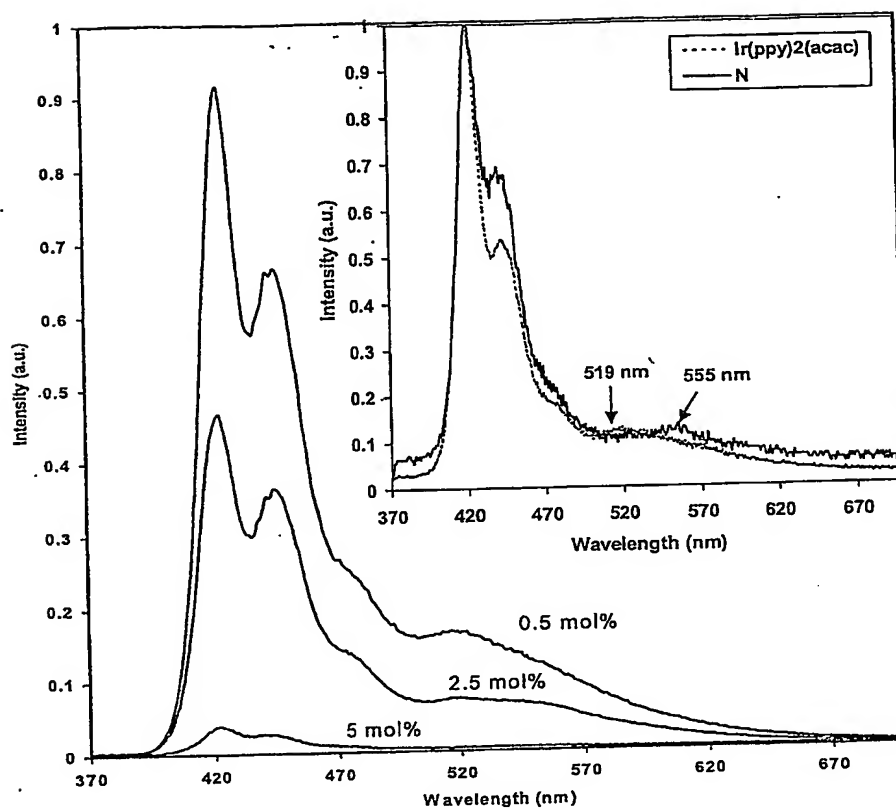
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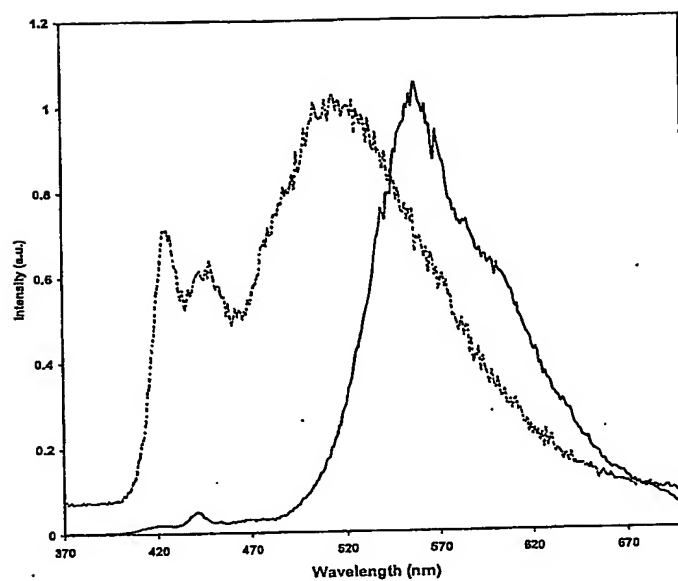
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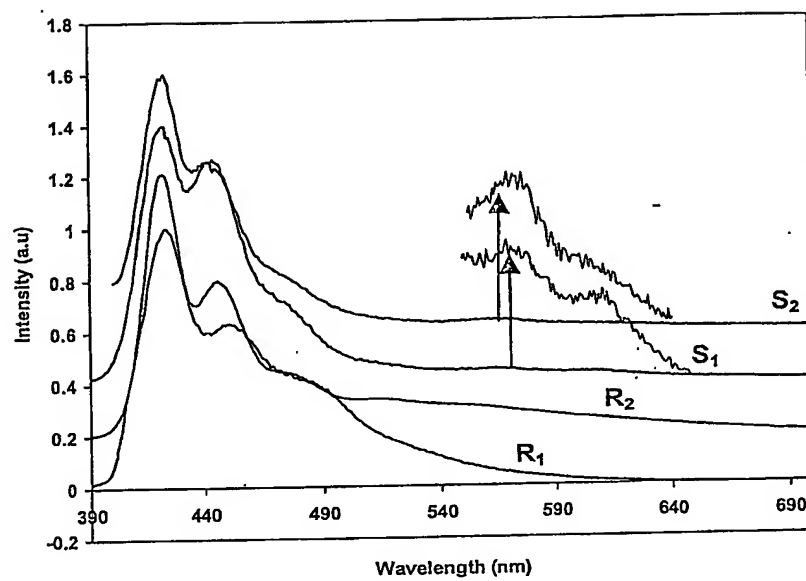
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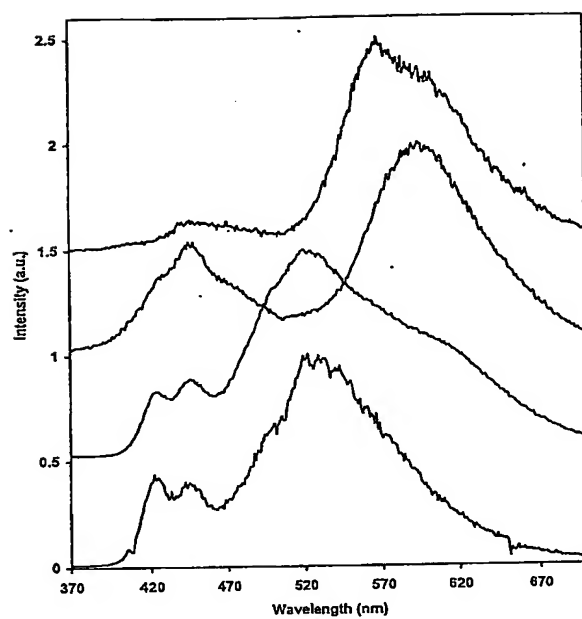
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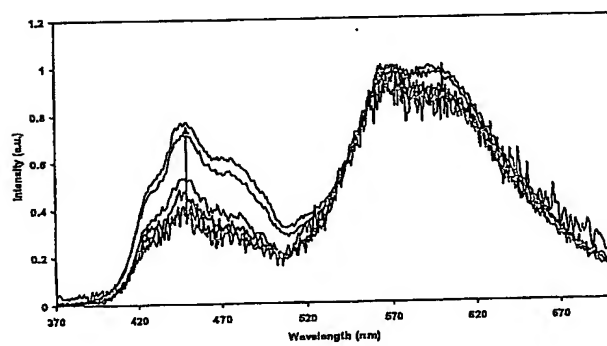
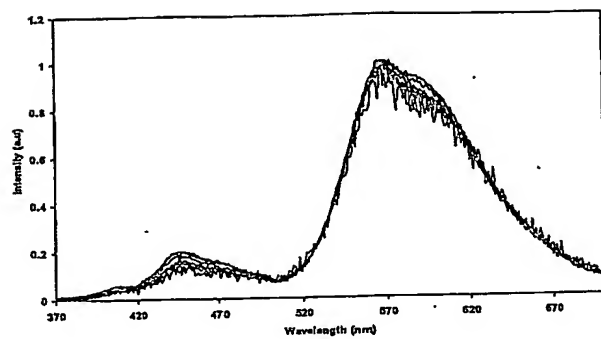
<Figure 5>



<Figure 6>



<Figure 7>



<Figure 8>

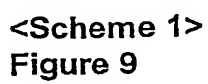
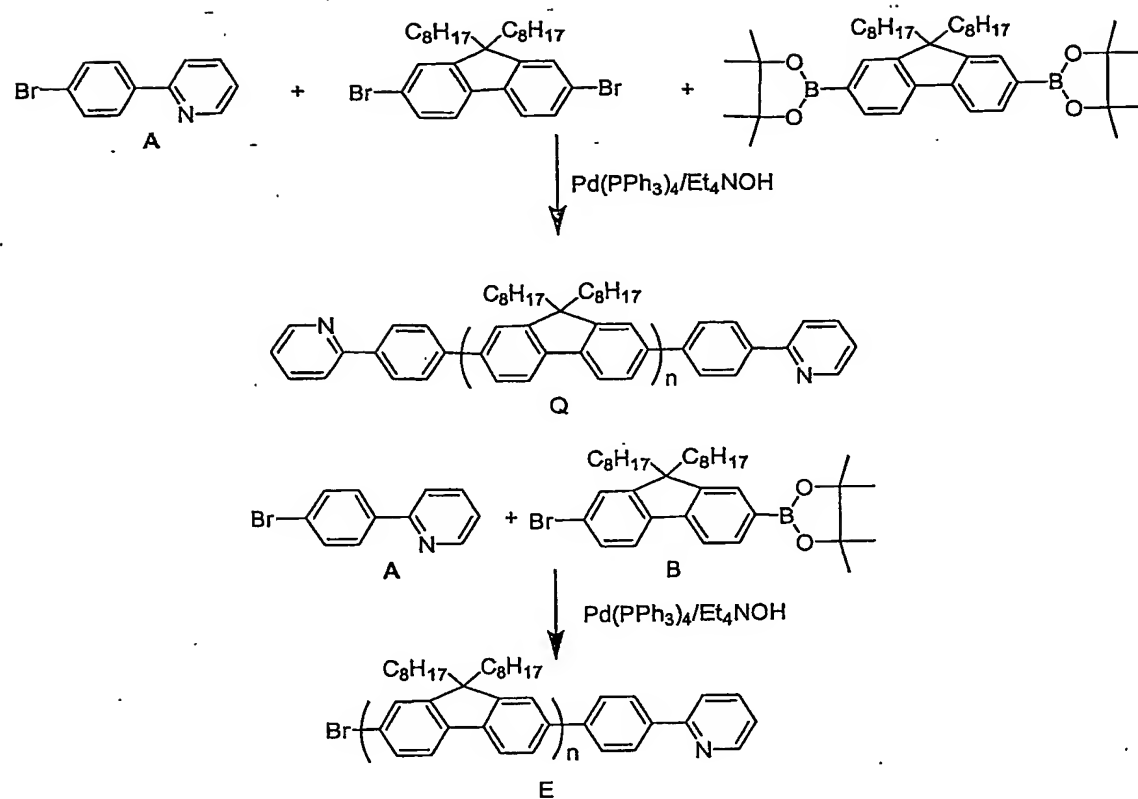
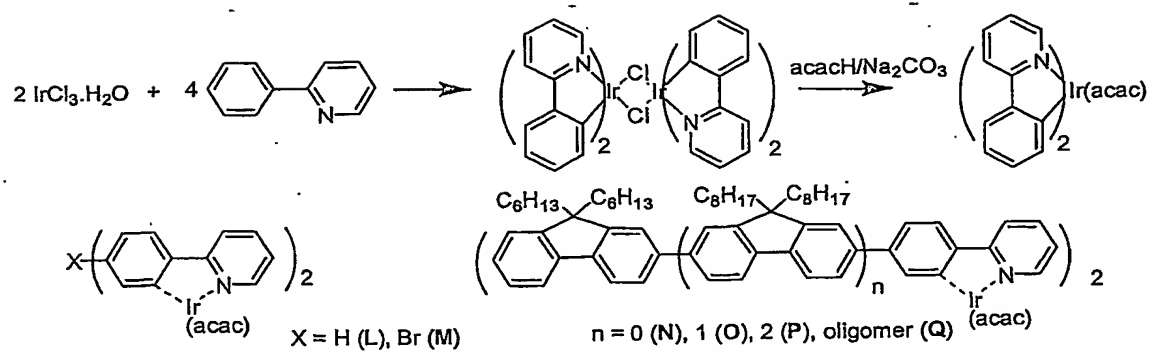


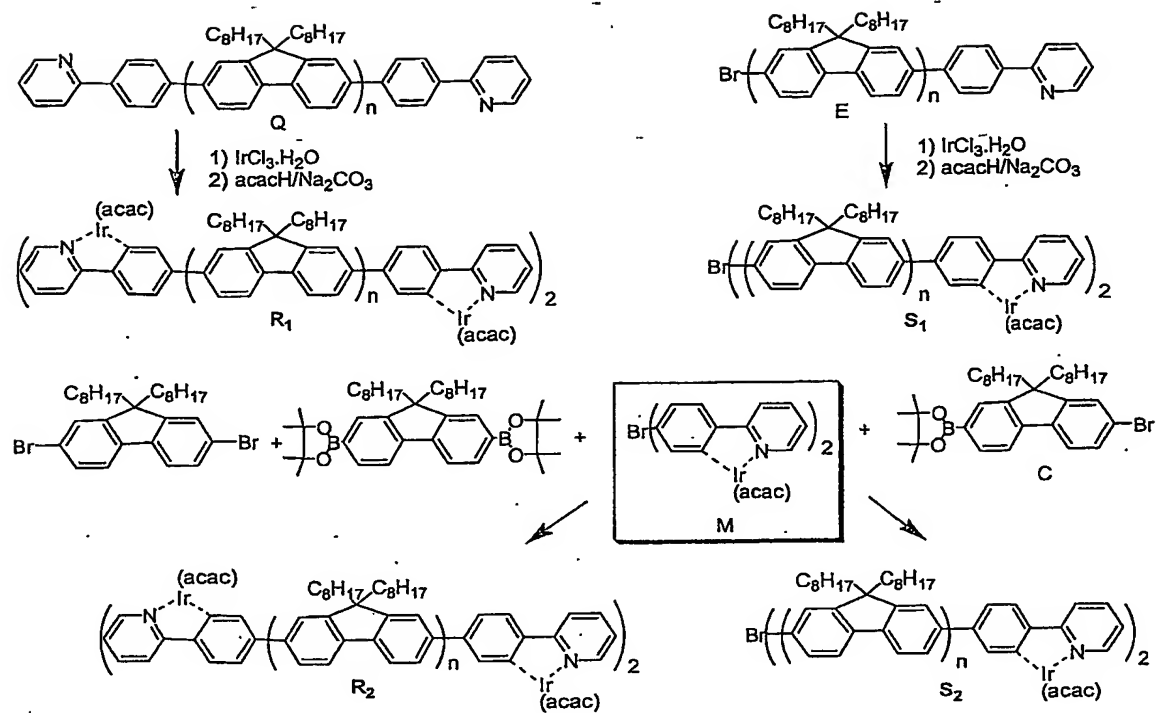
Figure 9



< Scheme 2 >
Figure 10



<Scheme 3>
Figure 11



<Scheme 4>
Figure 12

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